```
cathepsin.txt
? s cathepsin adj z or CTSZ
              0
                  CATHEPSIN ADJ Z
             60
                  CTSZ
S1
             60
                  S CATHEPSIN ADJ Z OR CTSZ
?
   rd
        Duplicate detection is not supported for File 393.
>>>W:
Duplicate detection is not supported for File 391.
Records from unsupported files will be retained in the RD set.
                  RD (UNIQUE ITEMS)
? t s29/free/1-29
>>>E: Set 29 does not exist
? t s29/3, k/1-29
>>>E: Set 29 does not exist
? t s1/3, k/1-29
>>>W: KWIC option is not available in file(s): 399
 1/3,K/1 (Item 1 from file: 5) Links
   Fulltext available through:
                                     USPTO Full Text Retrieval Options
Biosis Previews(R)
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0019601646 Biosis No.: 200700261387
Methylation markers common to breast and lung cancer segregate with breast cancer
risk in benign breast epithelial cells obtained by periareolar fine needle
aspiration (FNA).
Author: Euhus D M (Reprint); Shames D S; Lewis C M; Bu D; Minna J D
Author Address: UT SW Med Ctr, Dallas, TX USA**USA
Journal: Breast Cancer Research and Treatment 100 (Suppl. 1): p : Conference/Meeting: 29th Annual San Antonio Breast Cancer Symposium TX, USA December 14 -17, 2006; 20061214 Sponsor: San Antonio Canc Inst
                                                       100 ( Suppl. 1 ): p S226 2006 2006
                                                                              San Antonio,
Baylor Coll Med
Canc Therapy & Res Ctr
Univ Texas, Hlth Sci Ctr
ISSN: 0167-6806
Document Type: Meeting; Meeting Poster
Record Type: Citation
Language: English
DESCRIPTORS:
Gene Name: ...human CTSZ gene (Hominidae...
1/3,K/2 (Item 2 from file: 5) Links
Biosis Previews(R)
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18969215
            Biosis No.: 200600314610
Gene amplification in cancer
Author: Powers Scott; Yang Jianxin
Author Address: Greenlawn, NY USA**USA
Journal: Official Gazette of the United States Patent and Trademark Office Patents
DEC 13 2005 2005
Patent Number: US 06974672 Patent Date Granted: December 13. 2005 20051213 Patent
Classification: 435-6 Patent Assignee: Amgen Inc. Patent Country: USA
ISSN: 0098-1133
Document Type: Patent
Record Type: Abstract
Language: English
```

Abstract: ...diagnosis, prevention, and treatment of tumors and cancers in mammals, for example, humans, utilizing the CTSZ and CD24 genes, which are amplified colon cancer and/or ovarian cancer and/or breast cancer genes. The CTSZ and CD24 genes, their expressed protein products and antibodies are used diagnostically or as targets..

DESCRIPTORS:

Gene Name: ...human CTSZ gene (Hominidae)

1/3,K/3 (Item 3 from file: 5) Links

USPTO Full Text Retrieval Options Fulltext available through:

Biosis Previews(R)

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Biosis No.: 200600033869

Cathepsins in the ovine uterus: Regulation by pregnancy, progesterone, and

interferon tau

Author: Song Gwonhwa; Spencer Thomas E; Bazer Fuller W (Reprint)
Author Address: Texas A and M Univ, Ctr Anim Biotechnol and Genom, 442 Kleberg

Ctr,2471 TAMU, College Stn, TX 77843 USA**USA

Author E-mail Address: fbazer@cvm.tamu.edu

146 (11): p 4825-4833 NOV 2005 2005 Journal: Endocrinology

ISSN: 0013-7227

Document Type: Article Record Type: Abstract Language: English

Abstract: ...intracellular proteins, and processing of prohormones. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, and CTSZ genes was detected in the endometria of cyclic and early pregnant ewes with distinct temporal... expression patterns. In the d 18 and 20 conceptus, expression of CTSB, CTSD, CTSL, and CTSZ mRNA was detected in the trophectoderm. Of particular note, CTSL mRNA was the most abundant... ...tau. Other endometrial CTS genes were also regulated by progesterone alone (CTSB, CTSK, CTSS, and CTSZ) or progesterone and IFN tau (CTSH, CTSK, CTSS, and CTSZ). These results indicate that CTS of endometrial and conceptus origin may regulate endometrial remodeling and... DESCRIPTORS:

Gene Name: ...sheep CTSZ gene (Bovidae...

1/3,K/4 (Item 4 from file: 5) Links

Fulltext available through: USPTO Full Text Retrieval Options

Biosis Previews(R)

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18237633 Biosis No.: 200500144698

Cyclic tetrasaccharide-synthesizing enzymes from Arthrobacter globiformis A19

Author: Mukai Kazuhisa (Reprint); Maruta Kazuhiko; Satouchi Kazuhiro; Kubota Michio ; Fukuda Shigeharu; Kurimoto Masashi; Tsujisaka Yoshio

Author Address: Amase Inst, Hayashibara Biochem Labs Inc, 7-7 Amase Minami Machi,

Okayama, 7000834, Japan**Japan Author E-mail Address: amaseken@hayashibara.co.jp

Journal: Bioscience Biotechnology and Biochemistry 68 (12): p 2529-2540 December

2004 2004

Medium: print ISSN: 0916-8451

Document Type: Article Record Type: Abstract Language: English

Abstract: ...than the enzymes from strains of B. globisporus. The genes for IMT (ctsY) and 6GT (ctsZ) were cloned from the genome of A. globiformis A19. The two genes linked together in...

Page 2

DESCRIPTORS:

Gene Name: ...Arthrobacter globiformis ctsZ gene (Irregular Nonsporing Gram-Positive Rods)

1/3,K/5 (Item 5 from file: 5) Links

Fulltext available through: USPTO Full Text Retrieval Options

Biosis Previews(R)

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Biosis No.: 200200577451

Frequent amplification of 8q24, 11q, 17q, and 20q-specific genes in pancreatic cancer

Author: Mahlamaki Eija H (Reprint); Barlund Maarit; Tanner Minna; Gorunova Ludmila; Hoglund Mattias; Karhu Ritva; Kallioniemi Anne Author Address: Department of Clinical Chemistry, Tampere University Hospital,

FIN-33521, P.O. Box 2000, Tampere, Finland**Finland

35 (4): p 353-358 December, 2002 2002 Journal: Genes Chromosomes and Cancer

Medium: print ISSN: 1045-2257

Document Type: Article Record Type: Abstract Language: English

Abstract: ...17. In the 20q arm, the amplification frequencies varied from 32% to 83%, with the CTSZ gene at 20q13 being most frequently affected. These results illustrate that amplification of genes from...

1/3,K/6 (Item 6 from file: 5) Links

Fulltext available through: USPTO Full Text Retrieval Options

Biosis Previews(R)

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16867043 Biosis No.: 200200460554

Cloning and sequencing of the genes encoding cyclic tetrasaccharide-synthesizing enzymes from Bacillus globisporus C11

Author: Aga Hajime (Reprint); Maruta Kazuhiko; Yamamoto Takuo; Kubota Michio; Fukuda

Shigeharu; Kurimoto Masashi; Tsujisaka Yoshio

Author Address: Amase Institute, Hayashibara Biochemical Laboratories, 7-7 Amase

Minami-machi, Okayama, 700-0834, Japan**Japan

Journal: Bioscience Biotechnology and Biochemistry 66 (5): p 1057-1068 May, 2002 2002

Medium: print ISSN: 0916-8451

Document Type: Article Record Type: Abstract Language: English

Abstract: The genes for isomaltosyltransferase (CtsY) and 6-glucosyltransferase (CtsZ), involved in synthesis of a cyclic tetrasaccharide from alpha-glucan, have been cloned from the.....of 1093 residues having a signal sequence of 29 residues in its N-terminus. The ctsZ gene encodes a protein consisting of 1284 residues with a signal sequence of 35 residues...

1/3,K/7 (Item 7 from file: 5) Links

Fulltext available through: USPTO Full Text Retrieval Options

Biosis Previews(R)

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Biosis No.: 200000452781

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13

Author: Bonthron David T (Reprint); Hayward Bruce E; Moran Veronica; Strain Lisa Author Address: Molecular Medicine Unit, University of Leeds, St James's University

Hospital, Clinical Sciences Building, Leeds, LS9 7TF, UK**UK Journal: Human Genetics 107 (2): p 165-175 August, 2000 2000

Medium: print ISSN: 0340-6717

Document Type: Article Record Type: Abstract

Language: English

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20g13

Abstract: ...tissues, suggesting that, unlike GNAS1, TH1 is probably not imprinted. Immediately downstream of TH1 lies CTSZ, encoding the recently described cysteine protease, cathepsin Z. We have also elucidated the genomic structure.....TH1, only 70 bp separating their polyadenylation sites. A polymorphism was again identified within the CTSZ 3' untranslated region and used to demonstrate biallelic expression in fetal tissues.

DESCRIPTORS:

Chemicals & Biochemicals: ...human CTSZ gene...

1/3,K/8 (Item 1 from file: 24) Links
CSA Life Sciences Abstracts
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0002874280 IP Accession No: 6458347
Cyclic Tetrasaccharide-Synthesizing Enzymes from Arthrobacter globiformis A19

Mukai, Kazuhisa; Maruta, Kazuhiko; Satouchi, Kazuhiro; Kubota, Michio; Fukuda, Shigeharu; Kurimoto, Masashi; Tsujisaka, Yoshio Amase Institute, Hayashibara Biochemical Laboratories, Inc., 7-7 Amase minami-machi, Okayama 700-0834, Japan Bioscience, Biotechnology, and Biochemistry, v 68, n 12, p [np], 2004 Publication Date: 2004

Document Type: Journal Article

Record Type: Abstract Language: English

Summary Language: English

ISSN: 0916-8451

File Segment: Bacteriology Abstracts (Microbiology B)

Abstract:

...than the enzymes from strains of B. globisporus. The genes for IMT (ctsY) and 6GT (ctsZ) were cloned from the genome of A. globiformis A19. The two genes linked together in...

1/3, K/9 (Item 2 from file: 24) Links

Fulltext available through: USPTO Full Text Retrieval Options

CSA Life Sciences Abstracts

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0002450689 IP Accession No: 5569328

Cloning and Sequencing of the Genes Encoding Cyclic Tetrasaccharide-synthesizing Enzymes from Bacillus globisporus C11

Aga, Hajime; Maruta, Kazuhiko; Yamamoto, Takuo; Kubota, Michio; Fukuda, Shigeharu; Kurimoto, Masashi; Tsujisaka, Yoshio Amase Institute, Hayashibara Biochemical Laboratories, 7-7 Amase minami-machi, Okayama 700-0834, Japan, [mailto:amaseken@hayashibara.co.jp]

Bioscience, Biotechnology, and Biochemistry, v 66, n 5, p 1057-1068, May 2002

Publication Date: 2002

Document Type: Journal Article

Record Type: Abstract Language: English

Summary Language: English

ISSN: 0916-8451

File Segment: Bacteriology Abstracts (Microbiology B)

Abstract:

The genes for isomaltosyltransferase (CtsY) and 6-glucosyltransferase (CtsZ), involved in synthesis of a cyclic tetrasaccharide from alpha -glucan, have been cloned from the....of 1093 residues having a signal sequence of 29 residues in its N-terminus. The ctsZ gene encodes a protein consisting of 1284 residues with a signal sequence of 35 residues...

Descriptors: ...Aspartic acid; Gene clusters; cyclic tetrasaccharides; alpha -Glucan; 6-Glucosyltransferase; isomaltosyltransferase; glycoside hydrolase; ctsY gene; ctsZ gene; sugar transport; Bacillus globisporus; Thermococcus

1/3,K/10 (Item 1 from file: 34) Links
Fulltext available through: USPTO Full Text Retrieval Options
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14438813 Genuine Article#: 974IP No. References: 44
Cathepsins in the ovine uterus: Regulation by pregnancy, progesterone, and interferon tau

Author: Song GH; Spencer TE; Bazer FW (REPRINT) Corporate Source: Texas A&M Univ, Ctr Anim Biotechnol & Genom, 442 Kleberg Ctr, 2471 TAMU/College Stn//TX/77843 (REPRINT); Texas A&M Univ,Ctr Anim Biotechnol & Genom, College Stn//TX/77843; Texas A&M Univ, Dept Anim Sci, College Stn//TX/77843 (fbazer@cvm.tamu.edu) Journal: ENDOCRINOLOGY , 2005 , V 146 , N11 (NOV) , P 4825-4833 ISSN: 0013-7227 Publication date: 20051100 Publisher: ENDOCRINE SOC , 8401 CONNECTICUT AVE, SUITE 900, CHEVY CHASE, MD 20815-5817 USA Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE) Abstract: ...intracellular proteins, and processing of prohormones. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, and CTSZ genes was detected in the endometria of cyclic and early pregnant ewes with distinct temporal....expression patterns. In the d 18 and 20 conceptus, expression of CTSB, CTSD, CTSL, and CTSZ mRNA was detected in the trophectoderm. Of particular note, CTSL mRNA was the most abundant.....tau. Other endometrial CTS genes were also regulated by progesterone alone (CTSB, CTSK, CTSS, and CTSZ) or progesterone and IFN tau (CTSH, CTSK, CTSS, and CTSZ). These results indicate that CTS of endometrial and conceptus origin may regulate endometrial remodeling and...

1/3,K/11 (Item 2 from file: 34) Links
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13489190 Genuine Article#: 886TC No. References: 23
Cyclic tetrasaccharide-synthesizing enzymes from Arthrobacter globiformis A19
Author: Mukai K (REPRINT); Maruta K; Satouchi K; Kubota M; Fukuda S; Kurimoto M;
Tsujisaka Y
Corporate Source: Hayashibara Biochem Labs Inc,Amase Inst,7-7 Amase Minami
Machi/Okayama 7000834//Japan/ (REPRINT); Hayashibara Biochem Labs Inc,Amase
Inst,Okayama 7000834//Japan/ (amaseken@hayashibara.co.jp)
Journal: BIOSCIENCE BIOTECHNOLOGY AND BIOCHEMISTRY , 2004 , V 68 , N12 (DEC) , P
2529-2540

ISSN: 0916-8451 Publication date: 20041200 Page 5

cathepsin.txt Publisher: JAPAN SOC BIOSCI BIOTECHN AGROCHEM , JAPAN ACAD SOC CTR BLDG, 2-4-6 YAYOI BUNKYO-KU, TOKYO, 113, JAPAN Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)
Abstract: ...than the enzymes from strains of B. globisporus. The genes for IMT (ctsY) and 6GT (ctsZ) were cloned from the genome of A. globiformis A19. The two genes linked together in... 1/3,K/12 (Item 3 from file: 34) Links Fúlltext available through: USPTO Full Text Retrieval Options SciSearch(R) Cited Ref Sci (c) 2007 The Thomson Corp. All rights reserved. 11124549 Genuine Article#: 608ZV No. References: 23 Frequent amplification of 8q24, 11q, 17q, and 20q-specific genes in pancreatic cancer Author: Mahlamaki EH (REPRINT); Barlund M; Tanner M; Gorunova L; Hoglund M; Karhu R: Kallioniemi A Corporate Source: Tampere Univ Hosp, Dept Clin Chem, Canc Genet Lab, POB 2000/FIN-33521 Tampere//Finland/ (REPRINT); Tampere Univ Hosp, Dept Clin Chem, Canc Genet Lab, FIN-33521 Tampere//Finland/; Univ Tampere, FIN-33101 Tampere//Finland/; Univ Lund Hosp, Dept Clin Genet, S-22185 Lund//Sweden/
Journal: GENES CHROMOSOMES & CANCER, 2002, V 35, N4 (DEC), P 353-358
ISSN: 1045-2257 Publication date: 20021200 Publisher: WILEY-LISS , DIV JOHN WILEY & SONS INC, 605 THIRD AVE, NEW YORK, NY 10158-0012 USA Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE) Abstract: ...17. In the 20q arm, the amplification frequencies varied from 32% to 83%, with the CTSZ gene at 20q 13 being most frequently affected. These results illustrate that amplification of genes... 1/3,K/13 (Item 4 from file: 34) Links Fulltext available through: SciSearch(R) Cited Ref Sci USPTO Full Text Retrieval Options (c) 2007 The Thomson Corp. All rights reserved. 10692617 Genuine Article#: 557ZQ No. References: 33 10692617 Cloning and sequencing of the genes encoding cyclic tetrasaccharide-synthesizing enzymes from Bacillus globisporus C11 Author: Aga H (REPRINT); Maruta K; Yamamoto T; Kubota M; Fukuda S; Kurimoto M; Tsujisaka Y Corporate Source: Amase Inst, Hayashibara Biochem Labs, 77 Amase Minami Machi/Okayama 7000834//Japan/ (REPRINT); Amase Inst, Hayashibara Biochem Labs, Okayama 7000834//Japan/ Journal: BIOSCIENCE BIOTECHNOLOGY AND BIOCHEMISTRY, 2002, V 66, N5 (MAY), P 1057-1068 ISSN: 0916-8451 Publication date: 20020500 Publisher: JAPAN SOC BIOSCI BIOTECHN AGROCHEM, JAPAN ACAD SOC CTR BLDG, 2-4-6 YAYOI BUNKYO-KU, TOKYO, 113, JAPAN Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE) Abstract: The genes for isomaltosyltransferase (CtsY) and 6-glucosyltransferase (CtsZ), involved in synthesis of a cyclic tetrasaccharide from alpha-glucan, have been cloned from the.....of 1093 residues having a signal sequence of 29 residues in its N-terminus. The ctsZ gene encodes a protein consisting of 1284 residues with a signal sequence of 35 residues... 1/3,K/14 (Item 5 from file: 34) Links Fulltext available through: USPTO Full Text Retrieval Options SciSearch(R) Cited Ref Sci

No. References: 25

Page 6

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08981240

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13

Author: Bonthron DT (REPRINT); Hayward BE; Moran V; Strain L Corporate Source: UNIV LEEDS,ST JAMESS UNIV HOSP, MOL MED UNIT, CLIN SCI BLDG/LEEDS LS9 7TF/W YORKSHIRE/ENGLAND/ (REPRINT); UNIV EDINBURGH,WESTERN GEN HOSP, HUMAN GENET UNIT/EDINBURGH EH4 2XU/MIDLOTHIAN/SCOTLAND/ Journal: HUMAN GENETICS , 2000 , V 107 , N2 (AUG) , P 165-175 ISSN: 0340-6717 Publication date: 20000800 Publisher: SPRINGER-VERLAG , 175 FIFTH AVE, NEW YORK, NY 10010 Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE) Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13 Abstract: ...tissues, suggesting that, unlike GNAS1, TH1 is probably not imprinted. Immediately downstream of TH1 lies CTSZ, encoding the recently described cysteine protease, cathepsin Z. We have also elucidated the genomic structure... ...TH1, only 70 bp separating their polyadenylation sites. A polymorphism was again identified within the CTSZ 3' untranslated legion and used to demonstrate biallelic expression in fetal tissues.

1/3,K/15 (Item 1 from file: 50) Links
Fulltext available through: USPTO Full Text Retrieval Options
CAB Abstracts
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0008943406 CAB Accession Number: 20053216008
Cathepsins in the ovine uterus: regulation by pregnancy, progesterone, and interferon tau.

Song, G. H.; Spencer, T. E.; Bazer, F. W.
Author email address: fbazer@cvm.tamu.edu
Center for Animal Biotechnology and Genomics, Department of Animal Science, Texas
A&M University, College Station, TX 77843, USA.
Endocrinology vol. 146 (11): p.4825-4833
Publication Year: 2005
ISSN: 0013-7227
Digital Object Identifier: 10.1210/en.2005-0768
Publisher: Endocrine Society Bethesda , USA
Language: English Record Type: Abstract
Document Type: Journal article
... intracellular proteins, and processing of prohormones. Expression of CTSB, CTSD,
CTSH, CTSK, CTSL, CTSS, and CTSZ genes was detected in the endometria of cyclic and early pregnant ewes with distinct temporal..... expression patterns. In the d 18 and 20 conceptus, expression of CTSB, CTSD, CTSL, and CTSZ mRNA was detected in the trophectoderm. Of particular note, CTSL mRNA was the most abundant..... tau. Other endometrial CTS genes were also regulated by progesterone alone (CTSB, CTSS, and CTSZ) or progesterone and IFN (CTSH, CTSK, CTSS, and CTSZ). These results indicate that CTS of endometrial and conceptus origin may regulate endometrial

1/3,K/16 (Item 1 from file: 71) Links
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03839269 2007254436
Differential Expression of Cathepsins and Cystatin C in Ovine Uteroplacental Tissues

remodeling and ...

Song G.; Bazer F.W.; Spencer T.E. Address: T.E. Spencer, Center for Animal Biotechnology and Genomics, Department of Animal Science, Texas A and M University, 2471 TAMU, College Station, TX 77843-2471, United States

Email: tspencer@tamu.edu

Journal: Placenta, 28/10 (1091-1098), 2007, United Kingdom

CODEN: PLACD

ISSN: 0143-4004

Publisher Item Identifier: S0143400407001099

Document Type: Article

Summary Languages: English Languages: English

No. of References: 34

...roles in implantation and placentation in sheep. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, CTSZ, and CST3 mRNAs was detected in ovine uteroplacental tissues with distinct temporal and/or spatial expression patterns between Days 40 and 120 of pregnancy. Of particular note, CTSB, CTSD, and CTSZ mRNAs were predominantly detected in the chorion of the placenta and were more abundant in...

1/3, K/17 (Item 2 from file: 71) Links

Fulltext available through: USPTO Full Text Retrieval Options

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03106591

Cathepsins in the ovine uterus: Regulation by pregnancy, progesterone, and interferon tau

Song G.; Spencer T.E.; Bazer F.W.

Address: F.W. Bazer, Center for Animal Biotechnology and Genomics, 442 Kleberg Center, Texas A and M University, College Station, TX 77843-2471, United States

Email: fbazer@cvm.tamu.edu

Journal : Endocrinology , 146/11 (4825-4833) , 2005 , United States

CODEN: ENDOA ISSN: 0013-7227

Document Type: Article Languages: English

Summary Languages: English

No. of References: 44

...intracellular proteins, and processing of prohormones. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, and CTSZ genes was detected in the endometria of cyclic and early pregnant ewes with distinct temporal... ... expression patterns. In the d 18 and 20 conceptus, expression of CTSB, CTSD, CTSL, and CTSZ mRNA was detected in the trophectoderm. Of particular note, CTSL mRNA was the most abundant... ...tau. Other endometrial CTS genes were also regulated by progesterone alone (CTSB, CTSK, CTSS, and CTSZ) or progesterone and IFNtau (CTSH, CTSK, CTSS, and CTSZ). These results indicate that CTS of endometrial and conceptus origin may regulate endometrial remodeling and...

1/3,K/18 (Item 3 from file: 71) Links

Fulltext available through: USPTO Full Text Retrieval Options

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2005043458

Cyclic tetrasaccharide-synthesizing enzymes from Arthrobacter globiformis A19

Mukai K.; Maruta K.; Satouchi K.; Kubota M.; Fukuda S.; Kurimoto M.; Tsujisaka Y. Address: K. Mukai, Amase Institute, Hayashibara Biochem. Labs., Inc., 7-7 Amase

minami-machi, Okayama 700-0834 , Japan Email: amaseken@hayashibara.co.jp

Journal: Bioscience, Biotechnology and Biochemistry, 68/12 (2529-2540). 2004.

Japan

CODEN: BBBIE

ISSN: 0916-8451

Document Type: Article Languages: English Summary Languages: English

Page 8

No. of References: 23

...than the enzymes from strains of B. globisporus. The genes for IMT (ctsy) and 6GT (ctsZ) were cloned from the genome of A. globiformis A19. The two genes linked together in...

1/3,K/19 (Item 4 from file: 71) Links Fúlltext available through: USPTO Full Text Retrieval Options

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2004242374 02763829

Cloning and sequencing of the genes encoding cyclic tetrasaccharide- synthesizing enzymes from Bacillus globisporus C11

Aga H.; Maruta K.; Yamamoto T.; Kubota M.; Fukuda S.; Kurimoto M.; Tsujisaka Y. Address: H. Aga, Amase Institute, Hayashibara Biochemical Laboratories, 7-7 Amase minami-machi, Okayama 700-0834 , Japan

Email: amaseken@hayashibara.co.jp

Journal : Bioscience, Biotechnology and Biochemistry , 66/5 (1057-1068) , 2002 ,

Japan

CODEN: BBBIE ISSN: 0916-8451

Document Type: Article

Languages: English Summary Languages: English

No. of References: 33

The genes for isomaltosyltransferase (CtsY) and 6-glucosyltransferase (CtsZ), involved in synthesis of a cyclic tetrasaccharide from alpha-glucan, have been cloned from the.....of 1093 residues having a signal sequence of 29 residues in its N-terminus. The ctsZ gene encodes a protein consisting of 1284 residues with a signal sequence of 35 residues...

1/3,K/20 (Item 5 from file: 71) Links Fulltext available through: USPTO USPTO Full Text Retrieval Options **ELSEVIER BIOBASE**

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2002251489 02170500

Frequent amplification of 8q24, 11q, 17q, and 20q-specific genes in pancreatic cancer

Mahlamaki E.H.; Barlund M.; Tanner M.; Gorunova L.; Hoglund M.; Karhu R.; Kallioniemi A.

Address: E.H. Mahlamaki, Department of Clinical Chemistry, Tampere University Hospital, P.O. Box 2000, FIN-33521 Tampere , Finland

Email: eija.mahlamaki@tays.fi

Journal: Genes Chromosomes and Cancer, 35/4 (353-358), 2002, United States PUBLICATION DATE: December 1, 2002

CODEN: GCCAE ISSN: 1045-2257

Document Type: Article Languages: English No. of References: 23 Summary Languages: English

...17. In the 20q arm, the amplification frequencies varied from 32% to 83%, with the CTSZ gene at 20q 13 being most frequently affected. These results illustrate that amplification of genes...

1/3,K/21 (Item 6 from file: 71) Links Fulltext available through: USPTO Full Text Retrieval Options **ELSEVIER BIOBASE**

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01545037 2000208279

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13

Bonthron D.T.; Hayward B.E.; Moran V.; Strain L.

Address: D.T. Bonthron, Molecular Medicine Unit, University of Leeds, St. James's

University Hospital, Leeds LS9 7TF, United Kingdom

Email: D.T.Bonthron@leeds.ac.uk

Journal: Human Genetics, 107/2 (165-175), 2000, Germany

CODEN: HUGED ISSN: 0340-6717

Document Type: Article

Languages: English Summary Languages: English

No. of References: 26

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13

...tissues, suggesting that, unlike GNAS1, TH1 is probably not imprinted. Immediately downstream of TH1 lies CTSZ, encoding the recently described cysteine protease, cathepsin Z. We have also elucidated the genomic structure.....TH1, only 70 bp separating their polyadenylation sites. A polymorphism was again identified within the CTSZ 3' untranslated region and used to demonstrate biallelic expression in fetal tissues.

1/3,K/22 (Item 1 from file: 73) Links

Fulltext available through: USPTO Full Text Retrieval Options

EMBASE

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14746499 EMBASE No: 2007432049

Differential Expression of Cathepsins and Cystatin C in Ovine Uteroplacental Tissues

Song G.; Bazer F.W.; Spencer T.E. T.E. Spencer, Center for Animal Biotechnology and Genomics, Department of Animal Science, Texas A and M University, 2471 TAMU, College Station, TX 77843-2471 United States

Author Email: tspencer@tamu.edu

Placenta (PLACENTA) (United Kingdom) 2007 , 28/10 (1091-1098)

ISSN: 0143-4004 CODEN: PLACD

Publisher Item Identifier: S0143400407001099

Document Type: Journal; Article

Language: ENGLISH Summa Number Of References: 34 Summary Language: ENGLISH

...roles in implantation and placentation in sheep. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, CTSZ, and CST3 mRNAs was detected in ovine uteroplacental tissues with distinct temporal and/or spatial expression patterns between Days 40 and 120 of pregnancy. Of particular note, CTSB, CTSD, and CTSZ mRNAs were predominantly detected in the chorion of the placenta and were more abundant in...

1/3,K/23 (Item 2 from file: 73) Links

Fulltext available through: USPTO Full Text Retrieval Options

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EMBASE No: 2005463408

Cathepsins in the ovine uterus: Regulation by pregnancy, progesterone, and interferon tau

Song G.; Spencer T.E.; Bazer F.W. F.W. Bazer, Center for Animal Biotechnology and Genomics, 442 Kleberg Center, Texas Page 10

A and M University, College Station, TX 77843-2471 **United States**

Author Email: fbazer@cvm.tamu.edu

Endocrinology (ENDOCRINOLOGY) (United States) 2005 , 146/11 (4825-4833)

`ISSN: 0013-7227 CODEN: ENDOA Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 44

...intracellular proteins, and processing of prohormones. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, and CTSZ genes was detected in the endometria of cyclic and early pregnant ewes with distinct temporal....expression patterns. In the d 18 and 20 conceptus, expression of CTSB, CTSD, CTSL, and CTSZ mRNA was detected in the trophectoderm. Of particular note, CTSL mRNA was the most abundant....tau. Other endometrial CTS genes were also regulated by progesterone alone (CTSB, CTSK, CTSS, and CTSZ) or progesterone and IFNtau (CTSH, CTSK, CTSS, and CTSZ). These results indicate that CTS of endometrial and conceptus origin may regulate endometrial remodeling and...

1/3,K/24 (Item 3 from file: 73) Links

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EMBASE No: 2002389392 11819087

Frequent amplification of 8q24, 11q, 17q, and 20q-specific genes in pancreatic cancer

Mahlamaki E.H.; Barlund M.; Tanner M.; Gorunova L.; Hoglund M.; Karhu R.;

Kallioniemi A.

E.H. Mahlamaki, Department of Clinical Chemistry, Tampere University Hospital, P.O. Box 2000, FIN-33521 Tampere Finland

Author Email: eija.mahlamaki@tays.fi

Genes Chromosomes and Cancer (GENES CHROMOSOMES CANCER) (United States) 2002, 35/4 (353-358) CODEN: GCCAE ISSN: 1045-2257

Document Type: Journal; Article Language: ENGLISH Summary Langu

Summary Language: ENGLISH

Number of References: 23

...17. In the 20q arm, the amplification frequencies varied from 32% to 83%, with the CTSZ gene at 20q 13 being most frequently affected. These results illustrate that amplification of genes...

1/3,K/25 (Item 4 from file: 73) Links

Fulltext available through: USPTO Full Text Retrieval Options

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EMBASE No: 2000320402

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13

Bonthron D.T.; Hayward B.E.; Moran V.; Strain L. D.T. Bonthron, Molecular Medicine Unit, University of Leeds, St. James's University

Hospital, Leeds LS9 7TF United Kingdom

Author Email: D.T.Bonthron@leeds.ac.uk

Human Genetics (HUM. GENET.) (Germany) 2000 , 107/2 (165-175)

ISSN: 0340-6717 CODEN: HUGED

Document Type: Journal; Article

Summary Language: ENGLISH

Language: ENGLISH Summa Number Of References: 26

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20a13

Page 11

...tissues, suggesting that, unlike GNAS1, TH1 is probably not imprinted. Immediately downstream of TH1 lies CTSZ, encoding the recently described cysteine protease, cathepsin Z. We have also elucidated the genomic structure.....TH1, only 70 bp separating their polyadenylation sites. A polymorphism was again identified within the CTSZ 3' untranslated region and used to demonstrate biallelic expression in fetal tissues.

1/3,K/26 (Item 1 from file: 144) Links (c) 2007 INIST/CNRS. All rights reserved.

PASCAL No.: 05-0479921 17400127

Cathepsins in the ovine uterus : Regulation by pregnancy, progesterone, and interferon tau

GWONHWA SONG; SPENCER Thomas E; BAZER Fuller W Center for Animal Biotechnology and Genomics and Department of Animal Science, Texas A&M University, College Station, Texas 77843, United

Journal: Endocrinology: (Philadelphia),

2005, 146 (11) 4825-4833

Language: English

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... intracellular proteins, and processing of prohormones. Expression of CTSB, CTSD, CTSH, CTSK, CTSL, CTSS, and CTSZ genes was detected in the endometria of cyclic and early pregnant ewes with distinct temporal... expression patterns. In the d 18 and 20 conceptus, expression of CTSB, CTSD, CTSL, and CTSZ mRNA was detected in the trophectoderm. Of particular note, CTSL mRNA was the most abundant...

Other endometrial CTS genes were also regulated by progesterone alone (CTSB, CTSK, CTSS, and CTSZ) or progesterone and IFN tau (CTSH, CTSK, CTSS, and CTSZ). These results indicate that CTS of endometrial and conceptus origin may regulate endometrial remodeling and...

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PASCAL No.: 05-0144052 17078692

Cyclic tetrasaccharide-synthesizing enzymes from Arthrobacter globiformis A19

MUKAI Kazuhisa; MARUTA Kazuhiko; SATOUCHI Kazuhiro; KUBOTA Michio; FUKUDA Shigeharu; KURIMOTO Masashi; TSUJISAKA Yoshio Amase Institute, Hayashibara Biochemical Laboratories, Inc, 7-7 Amase minami-machi, Okayama 700-0834, Japan Journal: Bioscience, biotechnology, and biochemistry , 2004, 68 (12) 2529-2540 Language: English

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... than the enzymes from strains of B. globisporus. The genes for IMT (ctsY) and 6GT (ctsZ) were cloned from the genome of A. globiformis A19. The two genes linked together in...

1/3,K/28 (Item 3 from file: 144) Links Pascal (c) 2007 INIST/CNRS. All rights reserved.

15743060 PASCAL No.: 02-0454699

Cloning and sequencing of the genes encoding cyclic tetrasaccharide-synthesizing enzymes from Bacillus globisporus C11

AGA Hajime; MARUTA Kazuhiko; YAMAMOTO Takuo; KUBOTA Michio; FUKUDA Shigeharu; KURIMOTO Masashi; TSUJISAKA Yoshio Amase Institute, Hayashibara Biochemical Laboratories, 7-7 Amase minami-machi, Okayama 700-0834, Japan Journal: Bioscience, biotechnology, and biochemistry 2002, 66 (5) 1057-1068 Language: English

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The genes for isomaltosyltransferase (CtsY) and 6-glucosyltransferase (CtsZ), involved in synthesis of a cyclic tetrasaccharide from alpha -glucan, have been cloned from the...
... of 1093 residues having a signal sequence of 29 residues in its N-terminus. The ctsZ gene encodes a protein consisting of 1284 residues with a signal sequence of 35 residues...

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PASCAL No.: 01-0066263 14916379

Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13

BONTHRON David T: HAYWARD Bruce E; MORAN Veronica; STRAIN Lisa

Molecular Medicine Unit, University of Leeds, Clinical Sciences Building, St James's University Hospital, Leeds, LS9 7TF, United Kingdom; Human Genetics Unit, University of Edinburgh, Western General Hospital, Edinburgh, EH4 2XU, United Kingdom

Journal: Human genetics, 2000, 107 (2) 165-175

Language: English

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Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in chromosome 20q13 ...tissues, suggesting that, unlike GNASI, THI is probably not imprinted. Immediately downstream of THI lies CTSZ, encoding the recently described cysteine protease, cathepsin Z. We have also elucidated the genomic structure...

- ... TH1, only 70 bp separating their polyadenylation sites. A polymorphism was again identified within the CTSZ 3' untranslated region and used to demonstrate biallelic expression in fetal tissues.
- ...French Descriptors: Chromosome F20; Sequence nucleotide; Homologie; Organisation gene; Cathepsin; Origine parentale; Gene GNAS1; Gene TH1; Gene CTSZ

```
s cathepsin
S1
         88555
                   S CATHEPSIN
   s cathepsin(w)Z or CSTZ or cathepsin(w)X or cathepsin(w)P or cathepsin(w)Y
Processing
Processing
Processing
          88555
                   CATHEPSIN
       1099189
            152
                   CATHEPSIN(W)Z
             10
                   CSTZ
         88555
                   CATHEPSIN
       6940523
            197
                   CATHEPSIN(W)X
          88555
                   CATHEPSIN
      12346779
                   CATHEPSIN(W)P
             65
         88555
                   CATHEPSIN
       1989729
             30
                   CATHEPSIN(W)Y
S2
            450
                   S CATHEPSIN(W)Z OR CSTZ OR CATHEPSIN(W)X OR CATHEPSIN(W)P OR
CATHEPSIN(W)Y
   s s2 and compound
            450
                   S2
       4641049
                   COMPOUND
53
             67
                   S S2 AND COMPOUND
? rd
        Duplicate detection is not supported for File 393.
Duplicate detection is not supported for File 391.
Records from unsupported files will be retained in the RD set.
             63
                   RD (UNIQUE ITEMS)
? t s64/3.k/1-63
>>>E: Set 64 does not exist
? t s4/3, k/1-63
>>>W: KWIC option is not available in file(s): 399
 4/3,K/1 (Item 1 from file: 5) Links
   Fulltext available through:
                                      USPTO Full Text Retrieval Options
Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rights reserved.
           Biosis No.: 200510043834
18349334
Defining the substrate specificity of mouse cathepsin P
Author: Puzer Luciano; Barros Nilana M T; Oliveira Vitor; Julianoa Maria Aparecida;
Lu Guizhen; Hassanein Mohamed; Juliano Luiz; Mason Robert W; Carmona Adriana K
(Reprint)
Author Address: UNIFESP, Escola Paulista Med, Dept Biophys, Rua Tres Maio 100,
BR-04044020 Sao Paulo, Brazil**Brazil
Author E-mail Address: adriana@biofis.epm.br
Journal: Archives of Biochemistry and Biophysics
                                                              435 ( 1 ): p 190-196 MAR 1 05
2005
ISSN: 0003-9861
Document Type: Article
Record Type: Abstract
Language: English
Defining the substrate specificity of mouse cathepsin P
Abstract: Cathepsin P is a recently discovered placental cysteine protease that is structurally related to the more ubiquitously expressed, broad-specificity enzyme, cathepsin L. We studied the substrate specificity requirements of recombinant mouse
                                               Page 1
```

```
cathepsinsearch.txt
cathepsin P using fluorescence resonance energy transfer (FRET) peptides derived from the lead sequence Abz-KLRSSKQ-EDDnp......Arg), and hydrophobic aliphatic or aromatic residues (Val, Phe). For several substrates, the activity of cathepsin P was markedly regulated by kosmotropic salts, particularly Na2SO4. No significant effect on secondary or tertiary.....this substrate was almost two orders of
magnitude higher than that of the original parent compound. These results show that
cathepsin P, in contrast to other mammalian cathepsins, has a restricted catalytic
specificity. (C) 2004 Elsevier Inc...
Registry Numbers: ...cathepsin P
DESCRIPTORS:
 Chemicals & Biochemicals: ...cathepsin P--
  4/3,K/2 (Item 1 from file: 73)
                                              Links
     Fulltext available through:
                                              USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                  EMBASE No: 2007432049
14746499
Differential Expression of Cathepsins and Cystatin C in Ovine Uteroplacental Tissues
Song G.; Bazer F.W.; Spencer T.E. T.E. Spencer, Center for Animal Biotechnology and Genomics, Department of Animal Science, Texas A and M University, 2471 TAMU, College Station, TX 77843-2471
United States
Author Email: tspencer@tamu.edu
Placenta ( PLACENTA ) ( United Kingdom ) 2007 , 28/10 (1091-1098) CODEN: PLACD ISSN: 0143-4004
Publisher Item Identifier: S0143400407001099
Document Type: Journal; Article
Language: ENGLISH Summa
Number Of References: 34
                           Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cystatin C--endogenous compound--ec cathepsin B--endogenous compound--ec; cathepsin D--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin K--endogenous compound--ec;
cathepsin L--endogenous compound --ec; cathepsin S--endogenous compound--ec;
messenger RNA --endogenous compound--ec; peptide hydrolase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin Z-endogenous compound-ec
 4/3,K/3 (Item 2 from file: 73) Links
    Fulltext available through:
                                              USPTO Full Text Retrieval Options
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                 EMBASE No: 2007348195
14643786
Expression of Cathepsin P mRNA, Protein and Activity in the Rat Choriocarcinoma Cell
Line, Rcho-1, During Giant Cell Transformation
Hassanein M.; Korant B.D.; Lu G.; Mason R.W.
R.W. Mason, Department of Biomedical Research, Alfred I duPont Hospital for Children, 1600 Rock land Road, Wilmington, DE 19803 United States
Author Email: rmason@nemours.org
Placenta ( PLACENTA ) ( United Kingdom ) 2007 , 28/8-9 (912-919) CODEN: PLACD ISSN: 0143-4004 Publisher Item Identifier: S0143400406002773
Document Type: Journal; Article
Language: ENGLISH
                            Summary Language: ENGLISH
Number Of References: 44
Expression of Cathepsin P mRNA, Protein and Activity in the Rat Choriocarcinoma Cell
Line, Rcho-1, During Giant Cell...
...proteases perform critical functions in protein turnover and are essential for
```

Page 2

cathepsinsearch.txt normal growth and development. Cathepsin P is a member of a newly discovered family of lysosomal cysteine proteases uniquely expressed in... ... L was not regulated. A specific enzyme assay was developed to show that activity of cathepsin P mirrored mRNA expression during differentiation. Cathepsin P protein co-localizes with cathepsin B, indicating that the enzyme probably functions in the endosomal ... DRUG DESCRIPTORS: cathepsin--endogenous compound--ec cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec; peptide hydrolase--endogenous compound--ec; proteinase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin 1--endogenous compound--ec; cathepsin
2--endogenous compound--ec; cathepsin P--endogenous compound --ec; cathepsin
Q--endogenous compound--ec; cathepsin m--endogenous compound--ec; cathepsin r--endogenous compound--ec 4/3,K/4 (Item 3 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2007394692 Inflammatory processes in the aging mouse brain: Participation of dendritic cells and T-cells Stichel C.C.; Luebbert H. C.C. Stichel, Biofrontera Bioscience GmbH, D-51377 Leverkusen Germany Author Email: c.stichel-gunkel@biofrontera.com Neurobiology of Aging (NEUROBIOL. AGING) (United States) 2007 , 28/10 (1507-1521) ISSN: 0197-4580 CODEN: NEAGD Publisher Item Identifier: S0197458006002740 Document Type: Journal; Article Language: ENGLISH Summar Number Of References: 100 Summary Language: ENGLISH DRUG DESCRIPTORS: CD11b antigen--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin--endogenous compound--ec; integrin --endogenous compound--ec Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec 4/3,K/5 (Item 4 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. 14423842 EMBASE No: 2007158978 Differential expression of cathepsin X in aging and pathological central nervous system of mice Wendt W.; Zhu X.-R.; Lubbert H.; Stichel C.C. C.C. Stichel, Biofrontera Bioscience GmbH, D-51377 Leverkusen Germany Author Email: c.stichel-gunkel@biofrontera.com Experimental Neurology (EXP. NEUROL.) (United States) 2007, 204/2 (525-540) CODEN: EXNEA ISSN: 0014-4886 eISSN: 1090-2430 Publisher_Item Identifier: S0014488607000222 Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 77 Differential expression of cathepsin X in aging and pathological central nervous system of mice ...we analyzed the regional, cellular and subcellular localization and the activity of the recently discovered cathepsin X in the normal, developing and pathological mouse brain. Our results show that CATX is: (i... ...plaques in a transgenic mouse model and in Alzheimer patients. These results strongly suggest that cathepsin X is

Page 3

```
cathepsinsearch.txt
an important player in degenerative processes during normal aging and in pathological conditions. (c...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/6 (Item 5 from file: 73) Links
    Fulltext available through:
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                 EMBASE No: 2006599352
14198060
Cysteine cathepsins: Regulators of antitumour immune response
Obermajer N.; Doljak B.; Kos J.
J. Kos, University of Ljubljana, Department of Pharmaceutical Biology, Faculty of
Pharmacy, Askerceva 7, SI-1000 Ljubljana
                                                         Slovenia
Author Émail: Janko.kos@ffa.uni-lj.si
Expert Opinion on Biological Therapy (EXPERT OPIN. BIOL. THER. ) (United Kingdom)
   2006 , 6/12 (1295-1309)
CODEN: EOBTA ISSN: 1471-2598
Document Type: Journal; Review
Language: ENGLISH Summary Language: ENGLISH Number Of References: 120
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
cysteine derivative--endogenous compound--ec; cysteine proteinase --endogenous compound--ec; major histocompatibility antigen class 2 --endogenous compound--ec;
cytokine--endogenous compound--ec; growth factor--endogenous compound--ec; integrin--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin
L--endogenous compound--ec; cathepsin S--endogenous compound --ec; cathepsin K--endogenous compound--ec; stefin A--endogenous compound--ec; stefin B--endogenous compound--ec; cystatin C --endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin F--endogenous
compound--ec
Drug Terms (Uncontrolled): cathepsin w--endogenous compound--ec: cathepsin x
--endogenous compound--ec; cystatin f--endogenous compound --ec; cathepsin
o--endogenous compound--ec; cathepsin v--endogenous compound--ec
 4/3,K/7 (Item 6 from file: 73)
                                           Links
                                            USPTO Full Text Retrieval Options
    Fulltext available through:
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                EMBASE No: 2006493081
14090484
Cargo selectivity of the ERGIC-53/MCFD2 transport receptor complex
Nyfeler B.; Zhang B.; Ginsburg D.; Kaufman R.J.; Hauri H.-P.
H. Hans-Peter, Biozentrum, University of Basel, CH-4056 Basel Author Email: hans-peter.hauri@unibas.ch
                                                                                    Switzerland
Traffic (TRAFFIC) (Denmark) 2006, 7/11 (14 CODEN: TRAFF ISSN: 1398-9219 eISSN: 1600-0854 Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 37
                                                      7/11 (1473-1481)
...complex in the early secretory pathway. ERGIC-53 also interacts with the two
lysosomal glycoproteins cathepsin Z and cathepsin C. Here, we tested the subunit
interdependence and cargo selectivity of ERGIC-53... ... yellow fluorescent protein
fragment complementation. We found that MCFD2 is dispensable for the binding of
cathepsin Z and cathepsin C to ERGIC-53. The results indicate that ERGIC-53 can bind
cargo..
```

DRUG DESCRIPTORS:

```
cathepsinsearch.txt
* endoplasmic reticulum golgi intermediate compartment protein 53--endogenous compound--ec; *protein--endogenous compound--ec
secretory protein--endogenous compound--ec; receptor--endogenous compound--ec; blood clotting factor 5--endogenous compound --ec; blood clotting factor 8--endogenous
compound--ec; lectin --endogenous compound--ec; glycoprotein; dipeptidyl peptidase
I: cathepsin: protein subunit--endogenous compound--ec; small interfering RNA;
yellow fluorescent protein; unclassified drug
Drug Terms (Uncontrolled): multiple coagulation factor deficiency protein
2--endogenous compound--ec
 4/3,K/8 (Item 7 from file: 73)
                                        Links
    Fulltext available through:
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                EMBASE No: 2006471802
14057877
Cysteine cathepsins: Multifunctional enzymes in cancer
Mohamed M.M.; Sloane B.F.
B.F. Sloane, Department of Pharmacology, Wayne State University School of Medicine, Detroit, MI 48201 United States
Author Email: bsloane@med.wayne.edu
Nature Reviews Cancer ( NAT. REV. CANCER ) ( United Kingdom )
                                                                               2006 . 6/10
(764-775)
CODEN: NRCAC
                  ISSN: 1474-175X
Publisher Item Identifier: NRC1949
Document Type: Journal; Conference Paper
Language: ENGLISH
                        Summary Language: ENGLISH
Number Of References: 145
DRUG DESCRIPTORS:
* cysteine--endogenous compound--ec; *cathepsin--endogenous compound--ec cathepsin B--endogenous compound--ec; dipeptidyl peptidase I --endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin S--endogenous compound--ec; kininogen --endogenous compound--ec; cystatin C--endogenous compound
--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin O--endogenous compound--ec; cathepsin
V--endogenous compound--ec; cathepsin W--endogenous compound--ec; cathepsin
x--endogenous compound--ec; cystatin D --endogenous compound--ec; cystatin
E--endogenous compound --ec; cystatin f--endogenous compound--ec; cystatin
s--endogenous compound--ec; cystatin SA--endogenous compound--ec; cystatin
sn--endogenous compound--ec; cystatin M--endogenous compound --ec
 4/3,K/9 (Item 8 from file: 73)
                                        Links
    Fulltext available through:
                                        USPTO Full Text Retrieval Options
EMBASE
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                EMBASE No: 2006446396
Molecular aspects of stromal-parenchymal interactions in malignant neoplasms
Zalatnai A.
A. Zalatnai, First Department of Pathology and Experimental Cancer Research,
Semmelweis University, Faculty of Medicine, Ulloi ut 26, H-1085 Budapest Author Email: zalatnai@korb1.sote.hu
                                                                                            Hungary
Current Molecular Medicine ( CURR. MOL. MED. ) ( Netherlands )
                                                                                2006 , 6/6
(685 - 693)
                  ISSN: 1566-5240
CODEN: CMMUB
Document Type: Journal; Review
Language: ENGLISH Summai
Number Of References: 108
                       Summary Language: ENGLISH
DRUG DESCRIPTORS:
cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec; enzyme
                                                Page 5
```

cathepsinsearch.txt

precursor--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin K--endogenous compound--ec; coll47 antigen--endogenous compound--ec; interleukin lalpha--endogenous compound--ec; basic fibroblast growth factor--endogenous compound--ec; gelatinase B--endogenous compound--ec; tumor necrosis factor alpha--endogenous compound--ec; transforming growth factor beta--endogenous compound--ec; gelatinase A--endogenous compound--ec; collagenase 3--endogenous compound--ec; gelatinase A--endogenous compound--ec; stromal cell derived factor 1 --endogenous compound--ec; transforming growth factor betal --endogenous compound--ec; inducible nitric oxide synthase --endogenous compound--ec; gemcitabine--pharmacology--pd; matrix metalloproteinase inhibitor--drug therapy...

Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/10 (Item 9 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options

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13924109 EMBASE No: 2006345822

Caenorhabditis elegans: Study model for animal and human cathepsins and inhibitors

13924109 EMBASE No: 2006345822
Caenorhabditis elegans: Study model for animal and human cathepsins and inhibitors
Hashmi S.; Anwer K.; Bilgrami A.L.
S. Hashmi, Laboratory of Molecular Parasitology, Lindsley F. Kimbal Research
Institute, New York Blood Center, 310 East 67th Street, New York, NY 10021 United
States
Author Email: shashmi@nybloodcenter.org
Current Enzyme Inhibition (CURR. ENZYME INHIB.) (Netherlands) 2006 , 2/2
(173-188)
 ISSN: 1573-4080
Document Type: Journal; Review
Language: ENGLISH Summary Language: ENGLISH
Number Of References: 236

...L enzymes in C. elegans. Besides, it also reviews the function of a recently described cathepsin Z. (c) 2006 Bentham Science Publishers Ltd.

DRUG DESCRIPTORS:

* cathepsin-endogenous compound-ec; *cysteine proteinase inhibitor --drug therapy--dt; *cysteine proteinase inhibitor-endogenous compound-ec; *cysteine proteinase inhibitor--topical drug administration-tp cathepsin L-endogenous compound-ec; cathepsin B-endogenous compound-ec; cysteine proteinase-endogenous compound-ec; cathepsin F-endogenous compound-ec; cathepsin F-endogenous compound-ec; cathepsin F-endogenous compound-ec; cathepsin F-endogenous compound-ec; cathepsin K-endogenous compound-ec; cathepsin H-endogenous compound-ec; cathepsin K-endogenous compound-ec; stefin A-endogenous compound-ec; stefin B-endogenous compound-ec; anthelmintic agent; cyclophosphamide-drug therapy-dt; cyclophosphamide-opharmacology-pd; antineoplastic agent-drug therapy-dt; antineoplastic agent -pharmacology-pd; proteinase inhibitor-drug therapy-dt; proteinase inhibitor-endogenous compound-ec; proteinase inhibitor cystatin C-pharmacology-pd; proteinase inhibitor-topical drug administration-tp; antimalarial agent-opharmacology-pd; cystatin C-endogenous compound-ec; cystatin C-pharmacology-pd; cystatin C-endogenous compound-ec; cystatin C-pharmacology-pd; cystatin C-endogenous compound-ec; antivirus agent-opharmacology-pd; antivirus agent-opharmacology-pd; antivirus agent-opharmacology-pd; antivirus agent-opharmacology-pd; antivirus agent--topical drug administration-tp; unindexed drug; unclassified drug Drug Terms (Uncontrolled): cathepsin inhibitor--drug therapy-dt; cathepsin inhibitor--topical drug administration-ec; cathepsin inhibitor--pharmacology-pd; cathepsin inhibitor--topical drug administration-ec; cathepsin inhibitor--topical drug administration-ec; cathepsin inhibitor--topical drug administration-ec; cathepsin inhibitor--pharmacology-pd; alpha

ketoamide--drug therapy--dt...

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4/3,K/11 (Item 10 from file: 73) Links
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                     EMBASE No: 2006331563
Carboxypeptidase cathepsin X mediates betaSUB2-integrin-dependent adhesion of
differentiated U-937 cells
Obermajer N.; Premzl A.; Zavas(caron)nik Bergant T.; Turk B.; Kos J.
J. Kos, Faculty of Pharmacy, University of Ljubljana, As(carón)kerc(caron)eva 7, SI-1000 Ljubljana Slovenia
Author Email: janko.kos@ffa.uni-lj.si
Experimental Cell Research ( EXP. CELL RES. ) ( United States ) 01 AUG 2006 ,
312/13 (2515-2527)
                        ISSN: 0014-4827
CODEN: ECREA
Publisher Item Identifier: S0014482706001601
Document Type: Journal ; Article
                                Summary Language: ENGLISH
Language: ENGLISH
Number Of References: 50
Carboxypeptidase cathepsin X mediates betaSUB2-integrin-dependent adhesion of
differentiated U-937 cells
Cathepsin X is a lysosomal carboxypeptidase with a potential role in processes of
inflammation and immune response....integrin-binding motifs RGD and ECD, present in the pro- and in mature forms of cathepsin X, respectively, suggest that this
in the pro- and in mature forms of cathepsin X, respectively, suggest that this enzyme might have a function in cell signaling and adhesion. In.....protease inhibitors E-64 and CA-074 and 2F12 monoclonal antibody, all of which inhibit cathepsin X activity, significantly reduced adhesion of differentiated U-937 cells to polystyrene- and fibrinogen-coated surfaces.....whereas their binding to vitronectin, fibronectin or Matrigel was not affected. On the other hand, cathepsin X, added to differentiating U-937 cells, stimulated their adhesion. Using confocal microscopy, we demonstrated that the pro-form of cathepsin X was co-localized with betaSUB2 and betaSUB3 integrin subunits and its mature form solely with.....U-937 cells and in co-cultures with endothelial cells. Our results indicate that active cathepsin X mediates the function of betaSUB2 integrin receptors during cell adhesion and that it could also...
adhesion and that it could also...
DRUG DESCRIPTORS:
* carboxypeptidase--endogenous compound--ec; *beta2 integrin --endogenous
compound--ec
...cysteine proteinase inhibitor; monoclonal antibody; polystyrene; fibrinogen; integrin receptor; vitronectin; fibronectin; matrigel; beta3 integrin--endogenous
compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; ca 074
  4/3,K/12 (Item 11 from file: 73) Links
     Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                     EMBASE No: 2006270160
13831452
Tumor cell-derived and macrophage-derived cathepsin B promotes progression and lung
metastasis of mammary cancer
Vasiljeva O.; Papazoglou A.; Kruger A.; Brodoefel H.; Korovin M.; Deussing J.; Augustin N.; Nielsen B.S.; Almholt K.; Bogyo M.; Peters C.; Reinheckel T. T. Reinheckel, Institut fur Molekulare Medizin und Zellforschung,
Albert-Ludwigs-Universitat Freiburg, Stefan Meier Strasse 17, D-79104 Freiburg
Germany
Author Email: Thomas.Reinheckel@uniklinik-freiburg.de
Cancer Research ( CANCER RES. ) ( United States ) 15 MAY 2006 , 66/10 (5242-5250)
CODEN: CNREA ISSN: 0008-5472
```

Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH Number Of References: 48 ...labeling of cysteine cathepsins by the active site probe DCG-04 detected up-regulation of cathepsin X on PyMT;ctsbSUP+/+ cells. Treatment of cells with a neutralizing anti-cathepsin X antibody significantly reduced Matrigel invasion of PyMT;ctsbSUP+/+ cells but did not affect invasion of PyMT;ctsbSUP+/+ or PyMT;ctsbSUP+/- cells, indicating a compensatory function of cathepsin X in CTSB-deficient tumor cells. Finally, an adoptive transfer model, in which ctsbSUP+/+, ctsbSUP+/-, and... DRUG DESCRIPTORS: * cathepsin B--endogenous compound--ec virus middle T antigen--endogenous compound--ec; proteinase --endogenous compound--ec; matrigel--endogenous compound--ec; cysteine--endogenous compound--ec 4/3,K/13 (Item 12 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. 13797099 EMBASE No: 2006226285 13797099 Cysteine cathepsins in the immune response Zavas(caron)nik-Bergant T.; Turk B. B. Turk, Department of Biochemistry and Molecular Biology, J. Stefan Institute. SI-1000 Ljubljana Slovenia Author Email: boris.turk@ijs.si Tissue Antigens (TISSUE ANTIGENS) (United Kingdom) 2006 , 67/5 (349-355) ISSN: 0001-2815 eISSN: 1399-0039 CODEN: TSANA Document Type: Journal; Review Language: ENGLISH Summa Number Of References: 50 Summary Language: ENGLISH DRUG DESCRIPTORS: * cysteine derivative--endogenous compound--ec; *cathepsin--endogenous compound--ec major histocompatibility antigen class 2--endogenous compound--ec; cathepsin B--endogenous compound--ec; dipeptidyl peptidase I --endogenous compound--ec; cathepsin F--endogenous compound --ec; cathepsin H--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin D--endogenous compound --ec; CD4 antigen--endogenous compound--ec; CD8 antigen--endogenous compound--ec; major histocompatibility antigen class 1--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; cathepsin W--endogenous compound--ec; cathepsin V--endogenous compound --ec 4/3,K/14 (Item 13 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2004344716 13702260 Protease expression in interface tissues around loose arthroplasties Kido A.; Pap G.; Nagler D.K.; Ziomek E.; Menard R.; Neumann H.W.; Roessner A. Dr. A. Kido, Department of Orthopedic Surgery, Nara Medical University, 840 Shijo-cho, Kashihara, Nara 634-8522 Japan Author Email: akirakid@naramed-u.ac.jp Clinical Orthopaedics and Related Research (CLIN. ORTHOP. RELAT. RES.) (United States) 2004, -/425 (230-236) CODEN: CORTB ISSN: 0009-921X
Document Type: Journal; Article
Language: ENGLISH Summary Language
Number of Poff Summary Language: ENGLISH Number Of References: 35 Page 8

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DRUG DESCRIPTORS:
   proteinase--endogenous compound--ec
cathepsin--endogenous compound--ec; interstitial collagenase --endogenous
compound--ec; cathepsin B--endogenous compound --ec; cathepsin D--endogenous
compound--ec; cathepsin L--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
  4/3,K/15 (Item 14 from file: 73) Links
      Fulltext available through:
                                                             USPTO Full Text Retrieval Options
EMBASE
 (c) 2007 Elsevier B.V. All rights reserved.
                        EMBASE No: 2006135543
Lysosomal cysteine proteases: Structure, function and inhibition of cathepsins
Roberts R.
Prof. R. Roberts, Department of Biology, Program in Biochemistry and Molecular
Biology, Ursinus College, P.O. Box 1000, Collegeville, PA 19426-1000
                                                                                                                                    United States
Author Email: rroberts@ursinus.edu
Drug News and Perspectives ( DRUG NEWS PERSPECT. ) ( Spain )
                                                                                                                     2005 , 18/10
 (605-614)
                            ISSN: 0214-0934
CODEN: DNPEE
Document Type: Journal ; Review
Language: ENGLISH Summa
Number Of References: 111
                                   Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
lysosome enzyme--endogenous compound--ec; cysteine proteinase --endogenous
compound--ec; cathepsin B--endogenous compound --ec; dipeptidyl peptidase
I--endogenous compound--ec; cathepsin F --endogenous compound--ec; cathepsin
H--endogenous compound --ec; cathepsin F --endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin S--endogenous compound--ec; n [n (3 carboxyoxirane 2 carbonyl)leucyl]agmatine--drug comparison--cm; n [n (3... Drug Terms (Uncontrolled): cathepsin V--endogenous compound--ec; cathepsin O--endogenous compound--ec; cathepsin W--endogenous compound--ec; cathepsin w--endogenous compound--ec; cathepsin V--endogenous compound--ec; cathep
morpholineurea leucine homophenylalanine vinylsulfonephenyl--drug...
  4/3,K/16 (Item 15 from file: 73) Links
      Fulltext available through:
                                                             USPTO Full Text Retrieval Options
EMBASE
 (c) 2007 Elsevier B.V. All rights reserved.
                        EMBASE No: 2006036856
13558393
An enzyme-linked immunosorbent assay for human cathepsin X, a potential new
inflammatory marker
Nagler D.K.; Lechner A.M.; Oettl A.; Kozaczynska K.; Scheuber H.-P.;
Gippner-Steppert C.; Bogner V.; Biberthaler P.; Jochum M.
D.K. Nagler, Department of Clinical Chemistry and Clinical Biochemistry, University
Hospital of Surgery-City, Ludwig-Maximilians-University, Nussbaumstr. 20, 80336
                 Germany
Author Email: dorit.naegler@med.uni-muenchen.de
Journal of Immunological Methods ( J. IMMUNOL. METHODS ) ( Netherlands ) 2006 , 308/1-2 (241-250)
2006 , 308/1-
CODEN: JIMMB
                            ISSN: 0022-1759
Publisher Item Identifier: S0022175905003704
Document Type: Journal; Article
Language: ENGLISH
                                     Summary Language: ENGLISH
Number Of References: 35
An enzyme-linked immunosorbent assay for human cathepsin X, a potential new
inflammatory marker
                                                                         Page 9
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The human lysosomal cysteine-type carboxypeptidase cathepsin X is mainly present in
monocytes and macrophages and may be released into the circulation due.
 ...inflammatory marker, we have developed a highly sensitive and specific
sandwich-type immunoassay (ELISA) for cathepsin X permitting both intra- and extracellular detection and quantification. The dynamic range of the cathepsin X ELISA was determined to be 100 (detection limit) to 8000 pg/ml. Reproducibility of
both....of the thiol-dependent cathepsin family was not observed. The ELISA was used to quantify cathepsin X in leukocytes as well as in plasma of healthy volunteers and patients with multiple trauma. During the first 72 h after trauma, plasma levels of cathepsin X increased significantly, particularly in patients who died during the posttraumatic period. In comparison to the well-known inflammation marker neutrophil elastase, cathepsin X levels predicted survival with a higher significance in the later posttraumatic phase. In conclusion, this report provides the first evidence of cathepsin X immunoreactivity not only in cell lysates but also
in plasma samples. We suggest that the...
DRUG DESCRIPTORS:
 * cathepsin--endogenous compound--ec
cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec;
elastase--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
  4/3,K/17 (Item 16 from file: 73) Links
                                               USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
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                   EMBASE No: 2006015689
13530117
Endosomal proteases in antigen presentation
Chapman H.A.
H.A. Chapman, Department of Medicine, Cardiovascular Research Institute, University of California, San Francisco, CA 94143 United States
Author Email: hal.chapman@ucsf.edu
                                                                                                             2006,
Current Opinion in Immunology ( CURR. OPIN. IMMUNOL. ) ( United Kingdom )
18/1 (78-84)
CODEN: COPIE
                     ISSN: 0952-7915
Publisher Item Identifier: S0952791505002049
Document Type: Journal; Review
Language: ENGLISH Summ
Number Of References: 57
                             Summary Language: ENGLISH
DRUG DESCRIPTORS:
* proteinase--endogenous compound--ec; *major histocompatibility antigen class
1--endogenous compound--ec; *major histocompatibility antigen class 2--endogenous
compound--ec
cathepsin--endogenous compound--ec; cathepsin D--endogenous compound--ec; dipeptidyl
peptidase I--endogenous compound--ec;
cathepsin H--endogenous compound--ec; cathepsin S--endogenous compound--ec;
cathepsin L--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
  4/3,K/18 (Item 17 from file: 73) Links
     Fulltext available through: USPTO Full Text Retrieval Options
 (c) 2007 Elsevier B.V. All rights reserved.
                   EMBASE No: 2005489501
Gene expression profiles reveal increased mClca3 (Gob5) expression and mucin
production in a murine model of asbestos-induced fibrogenesis
Sabo-Attwood T.; Ramos-Nino M.; Bond J.; Butnor K.J.; Heintz N.; Gruber A.D.; Steele C.; Taatjes D.J.; Vacek P.; Mossman B.T. B.T. Mossman, University of Vermont, HSRF 218, 89 Beaumont Ave., Burlington, VT
                                                       Page 10
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United States
05405
Author Email: brooke.mossman@uvm.edu
American Journal of Pathology (AM. J. PATHOL.) (United States)
                                                                                                   2005 , 167/5
(1243 - 1256)
                     ISSN: 0002-9440
CODEN: AJPAA
Document Type: Journal; Article
Language: ENGLISH
                            Summary Language: ENGLISH
Number of References: 63
DRUG DESCRIPTORS:
* mucin--endogenous compound--ec; *asbestos; *gene product--endogenous compound--ec chrysotile; cyclin B1--endogenous compound--ec; cell cycle protein 20--endogenous
compound--ec; cyclin dependent kinase 1--endogenous compound--ec; chemokine--endogenous compound--ec; complement component C1--endogenous compound--ec; chitinase--endogenous compound--ec; tumor necrosis factor derivative--endogenous compound--ec; interleukin 1beta--endogenous compound--ec;
macrophage elastase--endogenous compound--ec; stromelysin --endogenous compound--ec;
integrin--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin
s--endogenous compound--ec; cytokine--endogenous compound--ec; unindexed drug;
unclassified drug
Drug Terms (Uncontrolled): protein mCLCA3--endogenous compound--ec; protein Gob5--endogenous compound--ec; CDC28 protein kinase regulatory subunit 2--endogenous
compound--ec; CCL9 chemokine--endogenous compound--ec; ccl6 chemokine--endogenous compound--ec; chitinase 3 like 3--endogenous compound--ec; tumor necrosis factor
superfamily member 10 --endogenous compound--ec; integrin alphax--endogenous
compound--ec; cathepsin Z--endogenous compound --ec
 4/3,K/19 (Item 18 from file: 73) Links
                                               USPTO Full Text Retrieval Options
     Fulltext available through:
EMBASE
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13365751 EMBASE No: 2005432915
Large scale real-time PCR analysis of mRNA abundance in rainbow trout eggs in relationship with egg quality and post-ovulatory ageing
Aegerter S.; Jalabert B.; Bobe J.
J. Bobe, INRA, SCRIBE, Campus de Beaulieu, F-35042 Rennes Cedex
                                                                                               France
Author Email: Julien.Bobe@rennes.inra.fr
Molecular Reproduction and Development ( MOL. REPROD. DEV. ) ( United States )
2005 , 72/3 (377-385)
CODEN: MREDE
                     ISSN: 1040-452X
Document Type: Journal ; Article
Language: ENGLISH Summary Language: ENGLISH Number Of References: 35
...period, eight transcripts (nucleoplasmin or Npm2, ferritin H, tubulin beta, JNK1,
cyclin A1, cyclin A2, cathepsin Z, and IGF2) exhibited a differential abundance at
one or several collection time(s). Interestingly, we....lower levels of Npm2, tubulin beta, and IGF1 transcripts. In contrast, keratins 8 and 18, cathepsin Z, and prostaglandin synthase 2 were more abundant in low quality eggs than in high
quality...
DRUG DESCRIPTORS:
nucleoplasmin--endogenous compound--ec; somatomedin--endogenous compound--ec; cyclin
A--endogenous compound--ec; tubulin --endogenous compound--ec; ferritin--endogenous compound--ec; stress activated protein kinase 1--endogenous compound--ec;
cathepsin--endogenous compound--ec; somatomedin C--endogenous compound--ec; beta tubulin--endogenous compound--ec; keratin --endogenous compound--ec; prostaglandin
synthase--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cyclin al--endogenous compound--ec; cyclin A2--endogenous
compound--ec; cathepsin Z--endogenous compound --ec; keratin 8--endogenous compound--ec; keratin 18--endogenous compound--ec; prostaglandin synthase 2--endogenous compound --ec
                                                       Page 11
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4/3,K/20 (Item 19 from file: 73) Links
    Fulltext available through:
                                          USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                 EMBASE No: 2005386832
Up-regulation of cathepsin X in Helicobacter pylori gastritis and gastric cancer
Krueger S.; Kalinski T.; Hundertmark T.; Wex T.; Kuster D.; Peitz U.; Ebert M.; Nagler D.K.; Kellner U.; Malfertheiner P.; Naumann M.; Rocken C.; Roessner A. S. Krueger, Department of Pathology, Otto-von-Guericke University, Leipziger Strasse 44, D-39120 Magdeburg Germany
Author Email: Sabine.Krueger@medizin.uni-magdeburg.de
Journal of Pathology ( J. PATHOL. ) ( United Kingdom ) CODEN: JPTLA ISSN: 0022-3417
                                                                         2005 , 207/1 (32-42)
CODEN: JPTLA
Document Type: Journal; Article
Language: ENGLISH
                         Summary Language: ENGLISH
Number Of References: 35
Up-regulation of cathepsin X in Helicobacter pylori gastritis and gastric cancer
Recently, we identified increased cathepsin X expression in H. pylori-infected gastric mucosa. Here, we describe further up-regulation in gastric cancer and report
on the role of inflammatory cytokines required for cathepsin X up-regulation in H.
pylori-infected gastric mucosa, as well as on consequences for cellular..
...infected and non-infected patients. Gastric cancer samples were obtained from
patients undergoing gastric surgery. Cathepsin X was detected in gastric mucosa by
quantitative real-time RT-PCR, western blotting and immunohistochemistry. Induction
of cathepsin X expression in epithelial and inflammatory cells caused by H. pylori infection was tested in in... ...cultures of AGS cells and monocytic cells. Patients
with H. pylori gastritis showed significantly higher cathepsin X mRNA (2.5-fold) and
protein (1.6-fold) expression than H. pylori-negative patients. Cathepsin X was also up-regulated in gastric cancer (3-12-fold) compared to non-neoplastic mucosa. Cathepsin X was predominantly expressed by macrophages in the mucosal stroma and in glands of the antral mucosa. In addition, tumour cells stained for cathepsin X in 26 (88%) patients with gastric carcinoma. In general, staining was significantly more
common (20... ... via soluble factors in the culture medium seems to be responsible
for increased expression of cathepsin X in monocytes. Using antisense
oligonucleotides, cathepsin X up-regulation was directly associated with higher
invasiveness in vitro. Although no correlation of cathepsin X expression and TNM
stage was found, our study demonstrates that cathepsin X plays a role not only in
the chronic inflammation of gastric mucosa but also in...
DRUG DESCRIPTORS:
  cathepsin--endogenous compound--ec
cytokine--endogenous compound--ec; messenger RNA--endogenous compound--ec; antisense
oligonucleotide; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/21 (Item 20 from file: 73) Links
    Fulltext available through:
                                          USPTO Full Text Retrieval Options
EMBASE
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                 EMBASE No: 2005224520
Carboxypeptidases cathepsins X and B display distinct protein profile in human cells
and tissues
Kos J.; Sekirnik A.; Premzl A.; Bergant V.Z.; Langerholc T.; Turk B.; Werle B.;
Golouh R.; Repnik U.; Jeras M.; Turk V.
J. Kos, Department of Pharmaceutical Biology, Faculty of Pharmacy, University of
Ljubljana, As(caron)kerc(caron)eva 7, SI-1000 Ljubljana
Author Email: janko.kos@krka.biz
Experimental Cell Research (EXP. CELL RES.) (United States)
                                                                                   15 MAY 2005 .
                                                 Page 12
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306/1 (103-113)

CODEN: ECREA ISSN: 0014-4827

Publisher Item Identifier: S0014482704007220

Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 42

Cathepsin X, a recently discovered lysosomal cysteine protease, shares common structural features and activity properties with cysteine.....distribution in cells and tissues and to their possible roles in malignancy. Protein level of cathepsin X did not differ significantly between matched pairs of lung tumor and adjacent lung tissue obtained......6-fold higher in tumor compared to adjacent lung tissue. Immunohistochemical analysis of lung tumor cathepsin X revealed very faint staining in tumor cells but positive staining in infiltrated histiocytes, alveolar macrophages, bronchial epithelial cells, and alveolar type II cells. Cathepsin X stained positive also in CD68SUP+ cells in germinal centers of secondary follicles in lymph nodes10A neoT and MDA-MB 231, showed positive staining for cathepsin B, but negative for cathepsin X. We showed that the invasive potential of MCF-10A neoT cells can be impaired by specific inhibitor of cathepsin B but not by that of cathepsin X. Cathepsin X was found in large amounts in the pro-monocytic U-937 cell line, in monocytes and in dendritic cells, generated from monocytes in vitro. Our results show that cathepsin X is not involved in degradation of extracellular matrix, a proteolytic event leading to tumor cell... DRUG DESCRIPTORS:

* carboxypeptidase--endogenous compound--ec; *cathepsin B--endogenous compound--ec CD68 antigen--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/22 (Item 21 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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13140612 EMBASE No: 2005207535
Capturing protein interactions in the secretory pathway of living cells

Nyfeler B.; Michnick S.W.; Hauri H.-P.
H.-P. Hauri, Dept. of Pharmacol. and Neurbiology, Biozentrum, University of Basel, Klingelbergstrasse 70, CH-4056 Basel Switzerland
Author Email: hans-peter.hauri@unibas.ch
Proceedings of the National Academy of Sciences of the United States of America (
PROC. NATL. ACAD. SCI. U. S. A.) (United States) 03 MAY 2005, 102/18
(6350-6355)

CODEN: PNASA ISSN: 0027-8424 Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 35

...ERGIC-53-interacting multicoagulation factor deficiency protein MCFD2, and to ERGIC-53's cargo glycoprotein cathepsin Z. YFP PCA analysis revealed the oligomerization of ERGIC-53 and its interaction with MCFD2, as well as its lectin-mediated interaction with cathepsin Z. Mutation of the lectin domain of ERGIC-53 selectively decreased YFP complementation with cathepsin Z. Using YFP PCA, we discovered a carbohydrate-mediated interaction between ERGIC-53 and cathepsin C...

DRUG DESCRIPTORS:

* endoplasmic reticulum golgi intermediate compartment protein 53--endogenous compound--ec; *cathepsin--endogenous compound--ec; *hybrid protein--endogenous compound--ec

Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; multicoagulation factor deficiency protein--endogenous compound--ec

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cathepsinsearch.txt
 4/3,K/23 (Item 22 from file: 73) Links
    Fulltext available through:
                                         USPTO Full Text Retrieval Options
EMBASE
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                EMBASE No: 2005198390
13138770
The human brain mannose 6-phosphate glycoproteome: A complex mixture composed of
multiple isoforms of many soluble lysosomal proteins
Sleat D.E.; Lackland H.; Wang Y.; Sohar I.; Xiao G.; Li H.; Lobel P.
Dr. P. Lobel, Ctr. for Adv. Biotech. and Medicine, 679 Hoes Lane, Piscataway, NJ
          United States
Author Email: lobel@cabm.rutgers.edu
Proteomics ( PROTEOMICS ,) ( Germany )
                                                   2005 , 5/6 (1520-1532)
CODEN: PROTC
                  ISSN: 1615-9853
Document Type: Journal ; Article
Language: ENGLISH
                         Summary Language: ENGLISH
Number Of References: 41
DRUG DESCRIPTORS:
  mannose 6 phosphate--endogenous compound--ec; *lysosome enzyme --endogenous
compound--ec; *proteome--endogenous compound--ec
somatomedin B receptor--endogenous compound--ec; n acetyl beta glucosaminidase--endogenous compound--ec; cathepsin S--endogenous compound--ec; deoxyribonuclease II--endogenous compound--ec; dipeptidyl peptidase--endogenous
compound--ec; gamma glutamyl hydrolase--endogenous compound--ec;
legumain--endogenous compound--ec; lysophospholipase--endogenous compound--ec;
proline carboxypeptidase--endogenous compound--ec; clusterin --endogenous
compound--ec; acetylesterase--endogenous compound--ec; alpha mannosidase--endogenous
compound--ec; serine carboxypeptidase--endogenous compound--ec; ribonuclease
--endogenous compound--ec; ependymin--endogenous compound--ec; ribonuclease
--endogenous compound--ec; ependymin--endogenous compound--ec; n4 (beta n
acetylglucosaminyl)asparaginase--endogenous compound --ec;
angiotensinogen--endogenous compound--ec; cerebroside sulfatase--endogenous
compound--ec; acylsphingosine deacylase --endogenous compound--ec; palmitoyl protein
thioesterase --endogenous compound--ec; cystatin C--endogenous compound --ec;
cystatin B--endogenous compound--ec; dipeptidyl peptidase I --endogenous
compound--ec; cathepsin D--endogenous compound --ec; cathepsin F--endogenous
compound--ec; cathepsin L--endogenous compound--ec; cathepsin--endogenous
compound--ec; F box protein--endogenous compound--ec; ferritin--endogenous
compound--ec; alpha levo fucosidase--endogenous compound--ec; alpha
glucosidase--endogenous compound--ec; sulfatase--endogenous compound--ec; alpha
galactosidase--endogenous compound--ec; beta galactosidase--endogenous compound--ec; beta glucuronidase --endogenous compound--ec; beta n acetylhexosaminidase
A--endogenous compound--ec; beta n acetylhexosaminidase B--endogenous compound--ec;
iduronate 2 sulfatase--endogenous compound--ec; levo iduronidase--endogenous compound--ec; galectin 1--endogenous compound--ec; acid lipase--endogenous
compound--ec; myelin associated glycoprotein--endogenous compound--ec; beta
mannosidase --endogenous compound--ec; prosaposin--endogenous compound --ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; cathepsin
p--endogenous compound--ec; cathepsin x --endogenous compound--ec; dipeptidyl
peptidase VII--endogenous compound--ec; ribonuclease 6--endogenous compound--ec; n
acetyl 6 galactosamine sulfatase--endogenous compound--ec; n acetyl glucosamine 6
sulfatase--endogenous compound--ec
 4/3,K/24 (Item 23 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
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                EMBASE No: 2005198358
13138755
Gene expression profiling of the effect of high-dose intravenous Ig in patients with
Kawasaki disease
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Abe J.; Jibiki T.; Noma S.; Nakajima T.; Saito H.; Terai M. Page 14

cathepsinsearch.txt Dr. J. Abe, Department of Allergy and Immunology, Natl. Res. Inst. Child Hlth./Devmt., 2-10-1 Ohkura, Setagaya-ku, Tokyo 157-8535 Author Email: jabe@nch.go.jp Journal of Immunology (J. IMMUNOL.) (United States) 01 MAY 2005 , 174/9 (5837 - 5845)CODEN: JOIMA ISSN: 0022-1767 Document Type: Journal; Article Language: ENGLISH Summa Number Of References: 53 Summary Language: ENGLISH DRUG DESCRIPTORS:

* ...dose--do; *immunoglobulin--drug therapy--dt; *immunoglobulin --intravenous drug administration--iv; *chemokine receptor CCR2--endogenous compound--ec; *protein S 100--endogenous compound--ec; *Fc receptor--endogenous compound--ec; *adrenomedullin--endogenous compound--ec formylpeptide receptor--endogenous compound--ec; C reactive protein --endogenous compound--ec; toll like receptor 2--endogenous compound--ec; adiponectin--endogenous compound--ec; cell surface receptor--endogenous compound--ec; colony stimulating factor receptor--endogenous compound--ec; interleukin 8 receptor --endogenous compound--ec; CD39 antigen--endogenous compound --ec; CD16 antigen--endogenous compound--ec; colony stimulating factor 1--endogenous compound--ec; protein tyrosine phosphatase --endogenous compound--ec; protein p57--endogenous compound --ec; interleukin 3--endogenous compound--ec; versican--endogenous compound--ec; immunoglobulin kappa chain--endogenous compound --ec; APRIL protein--endogenous compound--ec; dysferlin--endogenous compound--ec; chimerin--endogenous compound--ec; hematopoietic cell kinase--endogenous compound--ec; phosphatase --endogenous compound--ec; RGS2 protein--endogenous compound --ec; Rab protein--endogenous compound--ec; transcription factor -- endogenous compound--ec; protein v fos--endogenous compound --ec; early growth response factor 1--endogenous compound--ec; calreticulin--endogenous compound--ec; major histocompatibility antigen class 2--endogenous compound--ec; hexokinase--endogenous compound--ec; 5 aminolevulinate synthase--endogenous compound --ec; oxidoreductase--endogenous compound--ec; cytochrome P450 1B1 --endogenous compound--ec; long chain fatty acid coenzyme A ligase --endogenous compound--ec; histidine ammonialyase--endogenous compound--ec; microsomal aminopertidase--endogenous compound compound--ec; microsomal aminopeptidase--endogenous compound --ec; spermidine--endogenous compound--ec; acyltransferase --endogenous compound--ec; cathepsin--endogenous compound--ec; collapsin response mediator protein--endogenous compound--ec; ribosome protein--endogenous compound--ec; aquaporin 9--endogenous compound--ec; carrier protein--endogenous compound--ec; scramblase--endogenous compound--ec; heat shock protein 70 --endogenous compound--ec; protein disulfide isomerase--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): protein S100A9--endogenous compound--ec; protein
S100A12--endogenous compound--ec; protein S100A8--endogenous compound--ec;
adiponectin receptor 1--endogenous compound--ec; leukocyte immunoglobulin receptor B1--endogenous compound--ec; leukocyte immunoglobulin receptor B2--endogenous compound--ec; leukocyte immunoglobulin like receptor B3--endogenous compound--ec; stabilin 1 --endogenous compound--ec; S phase response protein--endogenous compound--ec; growth arrest specific protein 7--endogenous compound--ec; cold autoinflammatory syndrome 1 protein--endogenous compound--ec; pre B celĺ colony enhancing factor--endogenous compound--ec; proapoptotic caspase adaptor protein--endogenous compound--ec; chimerin 2--endogenous compound--ec; dual specificity phosphatase 1--endogenous compound--ec; Rab31 protein --endogenous compound--ec; kruppel like factor 4--endogenous compound--ec; cold shock domain protein A--endogenous compound--ec; SFFV proviral integration 1 protein--endogenous compound--ec; transcription factor 7 like 2--endogenous compound--ec; heads and accompound--ec; 3--endogenous compound--ec; guanosine phosphate reductase--endogenous compound--ec; biliverdin reductase B--endogenous compound--ec; gamma interferon inducible protein 30--endogenous compound--ec; flavoprotein oxidoreductase --endogenous compound--ec; neutrophil cytosolic factor 2--endogenous compound--ec; spermine n1 acetyltransferase--endogenous compound--ec; cathepsin Z--endogenous compound --ec;

mitochondrial solute carrier protein--endogenous compound--ec

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USPTO Full Text Retrieval Options
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EMBASE
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13086504 EMBASE No: 2005147588
Pancreatic beta-cell failure and diabetes in mice with a deletion mutation of the
endoplasmic reticulum molecular chaperone gene P58SUPIPK
Ladiges W.C.; Knoblaugh S.E.; Morton J.F.; Korth M.J.; Sopher B.L.; Baskin C.R.; MacAuley A.; Goodman A.G.; LeBoeuf R.C.; Katze M.G. W.C. Ladiges, Department of Comparative Medicine, Box 357190, University of
Washington, Seattle, WA 98195 United S
Author Email: wladiges@u.washington.edu
                                            United States
Diabetes ( DIABETES ) ( United States )
                                                          2005 , 54/4 (1074-1081)
CODEN: DIAEA ISSN: 0012-1797
Document Type: Journal; Article
Language: ENGLISH Summa
Number Of References: 26
                           Summary Language: ENGLISH
DRUG DESCRIPTORS:
* chaperone--endogenous compound--ec; *protein p58--endogenous compound--ec
glucose; initiation factor 2alpha-endogenous compound-ec; cathepsin L-endogenous compound-ec; protein p53-endogenous compound-ec; lymphotoxin beta-endogenous
compound--ec; cathepsin D--endogenous compound--ec; cathepsin B--endogenous
compound--ec; serine proteinase Omi--endogenous compound--ec; FAS ligand--endogenous
compound--ec; cathepsin--endogenous compound--ec; annexin--endogenous compound--ec;
cytochrome c'--endogenous compound--ec; STAT3 protein--endogenous compound --ec; beta arrestin--endogenous compound--ec; immunoglobulin enhancer binding
protein--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec
 4/3, K/26 (Item 25 from file: 73) Links
    Fulltext available through:
                                             USPTO Full Text Retrieval Options
(c) 2007 Elsevier B.V. All rights reserved.
                  EMBASE No: 2004261313
Cathepsins K, L, B, X and W are differentially expressed in normal and chronically
inflamed gastric mucosa
Buhling F.; Peitz U.; Kruger S.; Kuster D.; Vieth M.; Gebert I.; Roessner A.; Weber E.; Malfertheiner P.; Wex T. T. Wex, Dept. of Gastroenterology, Dept. of Infectious Disease, Leipziger Str. 44,
D-39120 Magdeburg Germany
Author Email: thomas.wex@medizin.uni-magdeburg.de
Biological Chemistry (BIOL. CHEM.) (Germany)
CODEN: BICHF ISSN: 1431-6730
                                                                       2004 , 385/5 (439-445)
Document Type: Journal; Article
Language: ENGLISH
                           Summary Language: ENGLISH
Number Of References: 28
...was expressed at very low levels. Infection by Helicobacter pylori caused a significant induction of cathepsin X (p<0.008), whereas the other cathepsins were
not or only locally affected by H. pylori infection or reflux disease.

Immunohistochemistry revealed specific expression of cathepsin X (macrophages),
cathepsin K (parietal cells) and cathepsin W (lymphocytes), whereas cathepsins B and
DRUG DESCRIPTORS:
* cathepsin K--endogenous compound--ec; *cathepsin L--endogenous compound--ec; *cathepsin B--endogenous compound--ec; * cathepsin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; cathepsin w
--endogenous compound--ec
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4/3, K/27 (Item 26 from file: 73) Links

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cathepsinsearch.txt
   Fulltext available through:
                                         USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
12657426
               EMBASE No: 2004255824
Up-regulation of cathepsin X in prostate cancer and prostatic intraepithelial
neoplasia
Nagler D.K.; Kruger S.; Kellner A.; Ziomek E.; Menard R.; Buhtz P.; Krams M.;
Roessner_A.; Keliner U.
D.K. Nagler, Dept. of Clin. Chem./Clin. Biochem., University Hospital of Surgery-City, Ludwig-Maximilians-University, Nussbaumstr. 20, 80336 Munich
                                                                                                Germany
Author Email: dorit.naegler@clinbio.med.uni-muenchen.de
Prostate ( PROSTATE ) ( United States ) 01 JUL 2004 , 60/2 (109-119) CODEN: PRSTD ISSN: 0270-4137
Document Type: Journal ; Article
Language: ENGLISH
                         Summary Language: ENGLISH
Number Of References: 36
Up-regulation of cathepsin X in prostate cancer and prostatic intraepithelial
neoplasia
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec
cathepsin F--endogenous compound--ec; cathepsin B--endogenous compound--ec;
cathepsin L--endogenous compound--ec; genomic DNA--endogenous compound--ec;
unclassified drug
prug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/28 (Item 27 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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                EMBASE No: 2004197020
12605998
Human B lymphoblastoid cells contain distinct patterns of cathepsin activity in
endocytic compartments and regulate MHC class II transport in a cathepsin
S-independent manner
Lautwein A.; Kraus M.; Reich M.; Burster T.; Brandenburg J.; Overkleeft H.S.;
Schwarz G.; Kammer W.; Weber E.; Kalbacher H.; Nordheim A.; Driessen C.
C. Driessen, MNF Universitat Tubingen, Ob dein Himmelreich 7, 72074 Tubingen
Author Email: christoph.driessen@med.uni-tuebingen.de
Journal of Leukocyte Biology ( J. LEUKOCYTE BIOL. ) ( United States )
                                                                                          2004 , 75/5
(844 - 855)
                  ISSN: 0741-5400
CODEN: JLBIE
Document Type: Journal; Article
                        Summary Language: ENGLISH
Language: ENGLISH
Number of References: 56
DRUG DESCRIPTORS:
* major histocompatibility antigen class 2--endogenous compound--ec; * cathepsin--endogenous compound--ec; *cathepsin S--endogenous compound--ec proteinase--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin D--endogenous compound--ec; dipeptidyl peptidase I--endogenous compound--ec; HLA DM antigen; vinyl derivative; phenol
derivative; unclassified drug
Drug Terms (Uncontrolled): asparagine specific endoprotease--endogenous
compound--ec; cathepsin Z; leucine homophenylalanine vinylsulfone phenol
 4/3, K/29 (Item 28 from file: 73) Links
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4/3,K/29 (Item 28 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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EMBASE No: 2004129918 12535984 Expression and characterization of cathepsin P Mason R.W.; Bergman C.A.; Lu G.; Frenck Holbrook J.; Sol-Church K. R.W. Mason, Department of Biomedical Research, Alfred I. duPont Hosp. for Children, 1600 Rockland Road, Wilmington, DE 19803 United States Author Email: mason@medsci.udel.edu Biochemical Journal (BIOCHEM. J.) (United Kingdom) 01 MAR 2004, 378/2 (657-663)CODEN: BIJOA ISSN: 0264-6021 Document Type: Journal; Article Language: ENGLISH Summa Number Of References: 24 Summary Language: ENGLISH Expression and characterization of cathepsin P ...in placental tissues of all mammalian species. In the present study, it was shown that cathepsin P can be expressed in Pichia pastoris as an inactive zymogen that can be activated with proteinase K, chymotrypsin or pancreatic elastase at neutral pH. Unlike other mammalian cathepsins, cathepsin P could also be autoactivated at neutral pH, but not at acidic pH. The activated enzyme.....SUB2SOSUB4 and hyaluronate stimulated the activity of the protease against peptidyl substrates. The properties of cathepsin P appear to be quite distinct from those of cathepsin L, indicating that the duplication that gave rise to cathepsin P has probably not yielded an enzyme that provides a subfunction of cathepsin L in rodents. It seems probable that cathepsin P has evolved to perform a function that is performed by an activities that cathepsin P has evolved to perform a function that is performed by an activities that cathepsin P has evolved to perform a function that is performed by an probable that cathepsin P has evolved to perform a function that is performed by an evolutionarily unrelated protease in... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cathepsin L--endogenous compound--ec messenger RNA--endogenous compound--ec; proteinase K; chymotrypsin; pancreatic elastase; peptide derivative--endogenous compound--ec; protein derivative--endogenous compound--ec; transferrin--endogenous compound--ec; inorganic salt; sodium sulfate; hyaluronic acid; unclassified drug Drug Terms (Uncontrolled): cathepsin p--endogenous compound--ec; azocasein --endogenous compound--ec 4/3,K/30 (Item 29 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2004127448 Cysteine proteases as disease markers Berdowska I. I. Berdowska, Department of Medical Biochemistry, Wroclaw Medical University, 10 Chalubinskiego, 50-368 Wroclaw Poland Author Email: iza@bioch.am.wroc.pl Clinica Chimica Acta (CLIN. CHIM. ACTA) (Netherlands) 2004, 342/1-2 (41-69) ISSN: 0009-8981 CODEN: CCATA Publisher Item Identifier: S0009898103006041 Document Type: Journal ; Review Language: ENGLISH Summa Number Of References: 248 Summary Language: ENGLISH DRUG DESCRIPTORS: * cysteine proteinase--endogenous compound--ec; *tumor marker --endogenous compound--ec peptidase--endogenous compound--ec; papain--endogenous compound--ec; cysteine derivative--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin

S--endogenous compound--ec; cathepsin K--endogenous compound --ec; cathepsin

F--endogenous compound--ec; dipeptidyl peptidase I --endogenous compound--ec; protein precursor--endogenous compound--ec; enzyme precursor--endogenous compound--ec; hormone precursor--endogenous compound--ec; major histocompatibility

Page 18

antigen class 2--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cysteine cathepsin derivative--endogenous compound--ec; cathepsin v -- endogenous compound--ec; cathepsin x--endogenous compound--ec; cathepsin w--endogenous compound--ec; cathepsin o--endogenous compound--ec 4/3.K/31 (Item 30 from file: 73) Links USPTO Full Text Retrieval Options Fulltext available through: **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. 12503843 EMBASE No: 2004098453 Myxobolus cerebralis: Identification of a cathepsin Z-like protease gene (MyxCP-1) expressed during parasite development in rainbow trout, Oncorhynchus mykiss Kelley G.O.; Adkison M.A.; Leutenegger C.M.; Hedrick R.P. G.O. Kelley, Dept. of Medicine and Epidemiology, School of Veterinary Medicine, University of California, Davis, CA 95616 United States Author Email: gokelley@ucdavis.edu Experimental Parasitology (EXP. PARASITOL.) (United States) 2003 , 105/3-4 (201-210)CODEN: EXPAA ISSN: 0014-4894 Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH
Number Of References: 54
Myxobolus cerebralis: Identification of a cathepsin Z-like protease gene (MyxCP-1) expressed during parasite development in rainbow trout, Oncorhynchus mykiss ...cysteine proteases. MyxCP-1 features a propeptide region and sequence insertions that are characteristics of cathepsin Z proteases. Phylogenetic comparisons of M. cerebralis to other eukaryotes based on full-length cathepsin-like genes show that MyxCP-1 is the earliest lineage in the cathepsin Z group and separated from cathepsin L, B, and C-like proteases. Using TaqMan PCR differential levels of transcription of the cathepsin Z-like protease were found in earlier and later developmental stages of the parasite in experimentally... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin Z like protease--endogenous compound-ec 4/3,K/32 (Item 31 from file: 73) Links USPTO Full Text Retrieval Options Fulltext available through: **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. 12487347 EMBASE No: 2004079961 The Caenorhabditis elegans Cathepsin Z-like Cysteine Protease, Ce-CPZ-1, Has a Multifunctional Role during the Worms' Development Hashmi S.; Zhang J.; Oksov Y.; Lustiqman S. S. Hashmi, Laboratory of Molecular Parasitology, Lindsley F. Kimball Research Inst., New York Blood Center, 310 E. 67th St., New York, NY 10021 United States Author Email: shashmi@nybloodcenter.org Journal of Biological Chemistry (J. BIOL. CHEM.) (United States) 279/7 (6035-6045)
CODEN: JBCHA ISSN: 0021-9258 13 FEB 2004, Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 60 The Caenorhabditis elegans Cathepsin Z-like Cysteine Protease, Ce-CPZ-1, Has a Multifunctional Role during the Worms' Development We have analyzed the expression and function of Ce-cpz-1, a Caenorhabditis elegans cathepsin Z-like cysteine protease gene, during development of the worm. The cpz-1

gene is expressed.....are degraded prior to shedding and ecdysis. The similar Page 19

localization of the related Onchocerca volvulus cathepsin Z protein suggests that the function of CPZ-1 during molting might be conserved in other... ...basement membrane extracellular matrix assembly process. The present findings have defined a critical role for cathepsin Z in nematode biology. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; cathepsin Z like cysteine proteinase--endogenous compound--ec

4/3,K/33 (Item 32 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2004009245 12409511 Identification of differentially expressed genes in models of melanoma progression by cDNA array analysis: SPARC, MIF and a novel cathepsin protease characterize aggressive phenotypes Rumpler G.; Becker B.; Hafner C.; McClelland M.; Stolz W.; Landthaler M.; Schmitt

R.; Bosserhoff A.; Vogt T. Dr. T. Vogt, Department of Dermatology, University of Regensburg, D-93042 Regensburg

Germany

Author Email: thomas.vogt@klinik.uni-regensburg.de

Experimental Dermatology (EXP. DERMATOL.) (United Kingdom) 2003 . 12/6 (761-771)

CODEN: EXDEE ISSN: 0906-6705 Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 46

...migration inhibiting factor (MIF), an important modulator of both cell cycle progression and angiogenesis, and cathepsin Z, a novel member of the family of matrix degrading proteinases. (c) Blackwell Munksgaard, 2003. DRUG DESCRIPTORS * complementary DNA--endogenous compound--ec; *osteonectin--endogenous compound--ec;

*macrophage migration inhibition factor--endogenous compound--ec; *cathepsin--endogenous compound--ec; * proteinase--endogenous compound--ec reduced nicotinamide adenine dinucleotide dehydrogenase (ubiquinone) --endogenous compound--ec; ubiquitin--endogenous compound--ec; selenoprotein--endogenous compound--ec; tumor protein--endogenous compound--ec; guanine nucleotide binding protein--endogenous compound--ec; HLA antigen class 2--endogenous compound--ec; laminin binding protein--endogenous compound--ec; protein --endogenous compound--ec; polyadenylic acid binding protein --endogenous compound--ec; DNA binding protein--endogenous compound--ec; initiation factor 2--endogenous compound--ec; heat shock protein 90--endogenous compound--ec; cytochrome b --endogenous compound--ec; protein lysine 6 oxidase--endogenous compound--ec; adenosine triphosphatase--endogenous compound --ec; receptor--endogenous compound--ec; phospholipid transfer protein--endogenous compound--ec; beta galactosidase--endogenous compound--ec; unindexed drug; unclassified drug Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; Wilm tumor related protein--endogenous compound--ec; guanine nucleotide binding protein beta subunit like protein--endogenous compound--ec; eukaryotic translation elongation factor 1 gamma--endogenous compound--ec; glia derived nexin--endogenous compound--ec; folic acid receptor 1--endogenous compound--ec

4/3,K/34 (Item 33 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2003402858 Phylogeny of antigen-processing enzymes: Cathepsins of a cephalochordate, an Page 20

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agnathan and a bony fish
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Uinuk-Ool T.S.; Takezaki N.; Kuroda N.; Figueroa F.; Sato A.; Samonte I.E.; Mayer
W.E.; Klein J.
T.S. Uinuk-Ool, Max-Planck-Inst. fur Biologie, Abteilung Immungenetik,
Corrensstrasse 42, D-72076 Tubingen
                                           Germany
Author Email: tanya@tuebingen.mpg.de
Scandinavian Journal of Immunology ( SCAND. J. IMMUNOL. ) ( United Kingdom )
                                                                                          01
OCT 2003, 58/4 (436-448)
CODEN: SJIMA
                 ISSN: 0300-9475
Document Type: Journal ; Article
Language: ENGLISH Summa
Number Of References: 71
                      Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec;
cathepsin F--endogenous compound--ec; dipeptidyl peptidase I--endogenous
compound--ec; cathepsin S --endogenous compound--ec; cathepsin K--endogenous
compound --ec; complementary DNA; unclassified drug
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; cathepsin O
--endogenous compound--ec
 4/3,K/35 (Item 34 from file: 73) Links
                                     USPTO Full Text Retrieval Options
   Fulltext available through:
EMBASE
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11901450
              EMBASE No: 2003014131
Determination of the mRNA sequence of cathepsin Y, a cysteine endopeptidase from rat
spleen, and confirmation of its ubiquitous expression
Nakazono E.; Kamata Y.; Yamafuji K.
K. Yamafuji, Division of Food and Nutrition, Nakamura Gakuen University, Befu 5-7-1,
Jonan-ku, Fukuoka 814-0198 Japan
Biological Chemistry (BIOL. CHEM.) (Germany)
CODEN: BICHF ISSN: 1431-6730
                                                          01 DEC 2002 , 383/12 (1971-1975)
Document Type: Journal; Article
Language: ENGLISH
                      Summary Language: ENGLISH
Number Of References: 14
Determination of the mRNA sequence of cathepsin Y, a cysteine endopeptidase from rat
spleen, and confirmation of its ubiquitous expression
...by its action of producing kinin-potentiating peptide from a plasma protein. We named it cathepsin Y due to its localization, acidic pH optimum and the presence of the same set of.....the mRNA sequence resulted in the omission of the strangely
attached C-terminal peptide from cathepsin Y.
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
messenger RNA--endogenous compound--ec; cysteine proteinase --endogenous
compound--ec; amino acid--endogenous compound --ec; thiol--endogenous compound--ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin Y--endogenous compound--ec
 4/3,K/36 (Item 35 from file: 73) Links
   Fulltext available through:
                                     USPTO Full Text Retrieval Options
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11309979
              EMBASE No: 2001324286
Lysosomal cysteine proteases: Facts and opportunities
Turk V.; Turk B.; Turk D. V. Turk, Department of Biochemistry, J. Stefan Institute, Ljubljana
                                                                               Slovenia
                                            Page 21
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Author Email: vito.turk@ijs.si
EMBO Journal ( EMBO J. ) ( United Kingdom )
                                                                03 SEP 2001 , 20/17 (4629-4633)
                     ISSN: 0261-4189
CODEN: EMJOD
Document Type: Journal; Review
                           Summary Language: ENGLISH
Language: ENGLISH
Number Of References: 45
DRUG DESCRIPTORS:
* cysteine proteinase--endogenous compound--ec; *cathepsin--endogenous compound--ec
enzyme precursor--endogenous compound--ec; amino acid; cathepsin L --endogenous
compound--ec; cathepsin S--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin H--endogenous compound--ec; dipeptidyl peptidase I--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin V--endogenous compound--ec; cathepsin w--endogenous compound--ec; cathepsin o--endogenous compound--ec; cathepsin
x--endogenous compound--ec
 4/3,K/37 (Item 36 from file: 73) Links
                                              USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                  EMBASE No: 2001244434
Cathepsins X and B display distinct activity profiles that can be exploited for
inhibitor design
Menard R.; Therrien C.; Lachance P.; Sulea T.; Qi H.; Alvarez-Hernandez A.; Roush
R. Menard, Biotechnology Research Institute, National Research Council of Canada,
6100 Royalmount Avenue, Montreal, Que. H4P 2R2
                                                                   Canada
Biological Chemistry (BIOL. CHEM.) (Germany)
CODEN: BICHF ISSN: 1431-6730
                                                                        2001 , 382/5 (839-845)
Document Type: Journal; Article
Language: ENGLISH Summary Language
Number Of References: 21
                           Summary Language: ENGLISH
  ..share similar activity profiles against substrates and inhibitors. Using quenched
fluorogenic substrates, we show that cathepsin X preferentially cleaves substrates
through a monopeptidyl carboxypeptidase pathway, while cathepsin B displays a
preference for.....approximately 2 orders of magnitude. Cleavage of a C-terminal
dipeptide of a substrate by cathepsin X can be observed under conditions that
preclude efficient monopeptidyl carboxypeptidase activity. In addition, an inhibitor designed to exploit the unique structural features responsible for the carboxypeptidase activity of cathepsin X has been synthesized and tested against cathepsins X, B and L. Although of moderate potency, this E-64 derivative is the first reported example of a cathepsin X-specific inhibitor. By comparison, CA074 was found to inactivate cathepsin B at least 34 000-fold more efficiently than cathepsin
DRUG DESCRIPTORS:
  cysteine proteinase--endogenous compound--ec; *cathepsin B --endogenous
compound--ec; *enzyme inhibitor--drug development--dv; *enzyme
inhibitor--pharmacology--pd
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; ca 074--drug
development--dv; ca 074--pharmacology--pd
 4/3,K/38 (Item 37 from file: 73) Links
    Fulltext available through:
                                              USPTO Full Text Retrieval Options
(c) 2007 Elsevier B.V. All rights reserved.
                  EMBASE No: 2001079403
Human cathepsin X: A novel cysteine protease with unique specificity
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cathepsinsearch.txt
Menard R.; Nagler D.K.; Zhang R.; Tam W.; Sulea T.; Purisima E.O. R. Menard, Biotechnology Research Institute, National Research Council of Canada, 6100 Avenue Royalmount, Montreal, Que. H4P 2R2 Canada
Advances in Experimental Medicine and Biology ( ADV. EXP. MED. BIOL. ) ( United
            2000 , 477/- (317-322)
CODEN: AEMBA ISSN: 0065-2598
Document Type: Journal : Conference Paper
Language: ENGLISH
Number Of References: 14
Human cathepsin X: A novel cysteine protease with unique specificity
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/39 (Item 38 from file: 73) Links
                                        USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
               EMBASE No: 2001079395
11062650
Review: Novel cysteine proteases of the papain family
Buhling F.; Fengler A.; Brandt W.; Welte T.; Ansorge S.; Nagler D.K. F. Buhling, Institue of Immunology, Otto von Guericke Univ. Magdeburg, Magdeburg
Germany
Advances in Experimental Medicine and Biology ( ADV. EXP. MED. BIOL. ) ( United
States ) 2000 , 477/- (241-254)
CODEN: AEMBA ISSN: 0065-2598
Document Type: Journal; Conference Paper
Language: ENGLISH
Number of References: 69
DRUG DESCRIPTORS:
* cysteine proteinase--endogenous compound--ec; *papain--endogenous compound--ec
cathepsin F--endogenous compound--ec; cathepsin K--endogenous compound--ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin o--endogenous compound--ec; cathepsin
V--endogenous compound--ec; cathepsin w--endogenous compound--ec; cathepsin
x--endogenous compound--ec
 4/3,K/40 (Item 39 from file: 73) Links
   Fulltext available through:
                                       USPTO Full Text Retrieval Options
EMBASE
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11023122
               EMBASE No: 2000123291
Proteolytic signals from Magdeburg
Ansorge S.; Langner J.; Buhling F.; Lendeckel U.
S. Ansorge, Inst. of Experimental Internal Med., Otto-von-Guericke University,
D-39120 Magdeburg Germany
Immunology Today ( IMMUNOL. TODAY ) ( United Kingdom ) 2000 , 21/4 (166-167)
CODEN: IMTOD ISSN: 0167-5699
Publisher Item Identifier: S0167569900015863
Document Type: Journal; Article
Language: ENGLISH
                        Summary Language: ENGLISH
DRUĞ DESCRIPTORS:
* microsomal aminopeptidase--endogenous compound--ec; *dipeptidyl peptidase
IV--endogenous compound--ec; *enzyme inhibitor--drug development--dv; *enzyme inhibitor--pharmacology--pd; *cathepsin --endogenous compound--ec
cathepsin S--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin D--endogenous compound--ec;
unclassified drug
                                               Page 23
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Drug Terms (Uncontrolled): peptidase inhibitor--drug development--dv; peptidase
inhibitor --pharmacology--pd; cathepsin w--endogenous compound--ec; cathepsin
F--endogenous compound--ec; cathepsin x--endogenous compound--ec
 4/3,K/41 (Item 40 from file: 73) Links
                                    USPTO Full Text Retrieval Options
   Fulltext available through:
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
10939353
              EMBASE No: 2000429272
Flow cytometric analysis of enzymes in live spermatozoa before and after cryostorage
Schaller J.; Glander H.-J.
Dr. J. Schaller, Dermatohistological Unit, Department of Dermatology, St. Barbara
Hospital, Barbarastr. 67, 47167 Duisburg
Andrologia (ANDROLOGIA) (Germany) 2
                                               Germany
                                            2000 . 32/6 (357-364)
                ISSN: 0303-4569
CODEN: ANDRD
Document Type: Journal ; Article
Language: ENGLISH
                      Summary Language: ENGLISH
Number Of References: 27
...for butyryl esterase (P<0.05), prolyl-aminopeptidase (P<0.001) and val-lys-(VK)-cathepsin (P<0.001) most probably due to elevated enzyme activities.
The activities of FDA-esterase (P...
DRUG DESCRIPTORS:
* peptidase--endogenous compound--ec; *proteinase--endogenous compound--ec; *esterase--endogenous compound--ec; *elastase --endogenous compound--ec;
*collagenase--endogenous compound --ec
fluorescein; rhodamine 110; microsomal aminopeptidase--endogenous compound--ec;
subtilisin--endogenous compound--ec; dipeptidyl peptidase--endogenous compound--ec;
proline iminopeptidase --endogenous compound--ec; cathepsin--endogenous compound--ec
 4/3,K/42 (Item 41 from file: 73) Links
   Fulltext available through:
                                    USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
              EMBASE No: 2000320402
Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in
chromosome 20q13
Bonthron D.T.; Hayward B.E.; Moran V.; Strain L. D.T. Bonthron, Molecular Medicine Unit, University of Leeds, St. James's University
Hospital, Leeds LS9 7TF
                           United Kingdom
Author Email: D.T.Bonthron@leeds.ac.uk
Human Genetics ( HUM. GENET. ) ( Germany )
                                                  2000 , 107/2 (165-175)
                ÌSSN: 0340-6717
CODEN: HUGED
Document Type: Journal; Article
Language: ENGLISH
                     Summary Language: ENGLISH
Number Of References: 26
...probably not imprinted. Immediately downstream of TH1 lies CTSZ, encoding the
recently described cysteine protease, cathepsin Z. We have also elucidated the
genomic structure of this gene; it has six exons spanning...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin z--endogenous compound--ec
 4/3,K/43 (Item 42 from file: 73) Links
   Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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Page 24

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cathepsinsearch.txt
(c) 2007 Elsevier B.V. All rights reserved.
                EMBASE No: 2000316209
10834786
Biochemical characterization of human cathepsin X revealed that the enzyme is an
exopeptidase, acting as carboxymonopeptidase or carboxydipeptidase
Klemencic I.; Carmona A.K.; Cezari M.H.S.; Juliano M.A.; Juliano L.; Guncar G.; Turk
D.; Krizaj I.; Turk V.; Turk B.
B. Turk, Dept. of Biochemistry/Molec. Biol., Josef Stefan Institute, Jamova 39, 1000
               Slovenia
Ljubljana
Author Email: boris.turk@ijs.si
European Journal of Biochemistry ( EUR. J. BIOCHEM. ) ( United Kingdom )
                                                                                              2000 .
267/17 (5404-5412)
CODEN: EJBCA
                   ISSN: 0014-2956
Document Type: Journal; Article
Language: ENGLISH Summa
Number Of References: 44
                         Summary Language: ENGLISH
Biochemical characterization of human cathepsin X revealed that the enzyme is an
exopeptidase, acting as carboxymonopeptidase or carboxydipeptidase
Cathepsin X, purified to homogeneity from human liver, is a single chain
glycoprotein with a molecular mass of approx. eq. 33 kDa and pI 5.1-5.3. Cathepsin X was inhibited by stefin A, cystatin C and chicken cystatin (K(i) = 1.7-15... ...was also inhibited by two specific synthetic cathepsin B inhibitors, CA-074 and GFG-semicarbazone. Cathepsin X was similar to cathepsin B and found to be a carboxypeptidase with preference for a positively charged Arg in P1 position.
Contrary to the preference of cathepsin B, cathepsin X normally acts as a
carboxymonopeptidase. However, the preference for Arg in the P1 position is so
strong that cathepsin X cleaves substrates with Arg in antepenultimate position,
acting also as a carboxydipeptidase. A large hydrophobic........P1' position,
although the enzyme cleaved all P1' residues investigated (Trp, Phe, Ala, Arg, Pro). Cathepsin X also cleaved substrates with amide-blocked C-terminal carboxyl group
with rates similar to those of the unblocked substrates. In contrast, no
endopeptidase activity of cathepsin X could be detected on a series of o-aminobenzoic acid-peptidyl-N-[2,-dinitrophenyl]ethylenediamine...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *carboxypeptidase--endogenous compound--ec;
*dipeptidase--endogenous compound--ec
liver enzyme--endogenous compound--ec; stefin A; cystatin C; stefin B; kininogen;
semicarbazone; cathepsin B; tryptophan; phenylalanine; arginine; proline...
 4/3,K/44 (Item 43 from file: 73) Links
                                         USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                EMBASE No: 2000138039
10728641
Role for cathepsin F in invariant chain processing and major histocompatibility
complex class II peptide loading by macrophages
Shi G.-P.; Bryant R.A.R.; Riese R.; Verhelst S.; Driessen C.; Li Z.; Bromme D.;
Ploegh H.L.; Chapman H.A.
H.A. Chapman, Pulmonary and Critical Care Div., University of California, 505
Parnassus Ave., San Francisco, CA 94143-0111
Author Email: halchap@itsa.ucsf.edu
                                                           United States
Journal of Experimental Medicine (J. EXP. MED.) (United States) 191/7 (1177-1185)
                                                                                        03 APR 2000 ,
```

...and dendritic cells revealed two enzymes expressed exclusively in macrophages, cathepsins Z and F. Recombinant cathepsin Z did not generate CLIP from Ii-MHC class Page 25

Summary Language: ENGLISH

CODEN: JEMEA

Language: ENGLISH

Number Of References: 42

ISSN: 0022-1007

Document Type: Journal; Article

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cathepsinsearch.txt
II complexes, whereas cathepsin F was as...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *major histocompatibility antigen class
2--endogenous compound--ec; *cell membrane protein --endogenous compound--ec
cysteine proteinase--endogenous compound--ec; recombinant enzyme; cathepsin S;
unclassified drug
Drug Terms (Uncontrolled): cathepsin F--endogenous compound--ec: class ii associated
invariant chain peptide--endogenous compound--ec: cathepsin z
 4/3,K/45 (Item 44 from file: 73) Links
   Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
10703930
              EMBASE No: 2000192563
The new subfamily of cathepsin-z-like protease genes includes Tc-cpz-1, a cysteine
protease gene expressed in Toxocara can's adults and infective stage larvae
Falcone F.H.; Tetteh K.K.A.; Hunt P.; Blaxter M.L.; Loukas A.; Maizels R.M.
R.M. Maizels, Inst. Cell Animal/Population Biol., University of Edinburgh, West
Mains Road, Edinburgh EH9 3JT
                                  United Kingdom
Author Email: r.maizels@ed.ac.uk
Experimental Parasitology (EXP. PARASITOL.) (United States)
                                                                      2000 , 94/3
(201-207)
                ISSN: 0014-4894
CODEN: EXPAA
Document Type: Journal ; Article
Language: ENGLISH
Number Of References: 32
The new subfamily of cathepsin-Z-like protease genes includes Tc-cpz-1, a cysteine
protease gene expressed in Toxocara canis...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin tc cpz 1--endogenous compound--ec
 4/3,K/46 (Item 45 from file: 73) Links
   Fulltext available through: USPTO Full Text Retrieval Options
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10651123
              EMBASE No: 2000116176
Mouse cathepsin M, a placenta-specific lysosomal cysteine protease related to
cathepsins L and P
Sol-Church K.; Frenck J.; Mason R.W.
R.W. Mason, Laboratory of Enzymology, Department of Research, Alfred I.du Pont Hosp.
for Children, P.O. Box 269, Wilmington, DE 19899 United States
Author Email: rmason@nemours.org
Biochimica et Biophysica Acta - Gene Structure and Expression ( BIOCHIM. BIOPHYS.
ACTA GENE STRUCT. EXPR. ) ( Netherlands ) 25 APR 2000 , 1491/1-3 (289-294)
               ISSN: 0167-4781
CODEN: BBGSD
Publisher Item Identifier: S0167478100000300
Document Type: Journal; Article
Language: ENGLISH Summa
Number Of References: 34
                     Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *complementary DNA--endogenous compound--ec;
*cysteine proteinase--endogenous compound--ec
cathepsin L--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin m--endogenous compound--ec; cathepsin p
```

4/3,K/47 (Item 46 from file: 73) Links

--endogenous compound--ec

Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000116156 10651103 Murine and human cathepsin Z: cDNA-cloning, characterization of the genes and chromosomal localization Deussing J.; Von Olshausen I.; Peters C. C. Peters, Institut Molekular Medizin, Klinikum, Albert-Ludwig-Universitat, Hugstetter Strasse 55, 7910b Freiburg German, Author Email: peters@mm11.ukl.uni-freiburg.de
Biochimica et Biophysica Acta - Gene Structure and Expression (BIOCHIM. BIOPHYS.

ACTA GENE STRUCT. EXPR.) (Netherlands)
CODEN: BBGSD ISSN: 0167-4781
Publisher Item Identifier: S016747810000021X

Document Type: Journal ; Article

Summary Language: ENGLISH Language: ENGLISH

Number Of References: 53

Murine and human cathepsin Z: cDNA-cloning, characterization of the genes and chromosomal localization

...encoding a predicted polypeptide of 306 amino acids was characterized. The new protease, tentatively named cathepsin Z, exhibits all features characteristics of a papain-like cysteine protease, including the highly conserved residues of the 'catalytic triad'. Cathepsin Z shares only 26-35% overall homology with previously described mammalian papain-like cysteine peptidases and.....within the family of papain-like cysteine peptidases. Genomic clones covering the murine and human cathepsin Z genes were isolated. They comprise six exons and five introns spanning a 12-kb region of genomic DNA, respectively. Murine cathepsin Z was mapped to chromosome 2, a region with synteny homology to a region of human chromosome 20 to which human cathepsin Z has been mapped previously. Northern blot analysis revealed ubiquitous expression of murine cathepsin Z. Multiple transcriptional start sites were identified for the murine cathepsin Z gene and together with the absence of a TATA box, a high G+C content.....CpG island and the presence of several Sp1-binding sites in the promoter region, murine cathepsin Z may be classified as a 'housekeeping' gene. Copyright (C) 2000 Elsevier Science B.V. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec complementary DNA--endogenous compound--ec

4/3,K/48 (Item 47 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000099482 10633942 Crystal structure of cathepsin X: A flip-flop of the ring of His23 allows carboxy-monopeptidase and carboxy-dipeptidase activity of the protease

Guncar G.; Klemencic I.; Turk B.; Turk V.; Karaoglanovic-Carmona A.; Juliano L.; Turk D.

D. Turk, Dept. of Biochem./Molecular Biology, Jozef Stefan Institute, Jamova 39, 1000 Ljubljana Slovenia

Author Email: dusan.turk@ijs.si

15 MAR 2000 , 8/3 (305-313)

Structure (STRUCTURE) (United Kingdom)
CODEN: STRUE ISSN: 0969-2126 Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 54

Crystal structure of cathepsin X: A flip-flop of the ring of His23 allows carboxy-monopeptidase and carboxy-dipeptidase activity...

Background: Cathepsin X is a widespread, abundantly expressed papain- like mammalian Page 27

lysosomal cysteine protease. It exhibits carboxy-monopeptidase.....of the two enzyme activities has actually been monitored. Results: The crystal structure of human cathepsin X has been determined at 2.67 Angstrom resolution. The structure shares the common features of.....like enzyme fold, but with a unique active site. The most pronounced feature of the cathepsin X structure is the mini-loop that includes a short three- residue insertion protruding into the... ...terminal carboxyl group of a substrate in two different sidechain conformations. Conclusions: The structure of cathepsin X exhibits a binding surface that will assist in the design of specific inhibitors of cathepsin X as well as of cathepsin B and thereby help to clarify the physiological roles of... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *histidine--endogenous compound--ec; *peptidase--endogenous compound--ec; * proteinase--endogenous compound--ec 4/3,K/49 (Item 48 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000067209 10601951 Cathepsin Q, a novel lysosomal cysteine protease highly expressed in placenta Sol-Church K.; Frenck J.; Mason R.W. R.W. Mason, Laboratory of Enzymology, Department of Research, Alfred I. duPont Hospital Children, PO Box 269, Wilmington, DE 19899 United States Author Email: rmason@nemours.org Biochemical and Biophysical Research Communications (BIOCHEM. BIOPHYS. RES. COMMUN.) (United States) 27 JAN 2000 , 267/3 (791-795) ISSN: 0006-291X CODEN: BBRCA Document Type: Journal ; Article Language: ENGLISH Summa Number Of References: 22 Summary Language: ENGLISH ...is predicted that cathepsin Q will differ in catalytic specificity to another placental-specific protease, cathepsin P, indicating that these enzymes will have unique proteolytic functions in extra-embryonic tissues. (C) 2000... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec; *lysosome enzyme--endogenous compound--ec 4/3,K/50 (Item 49 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. 10531360 EMBASE No: 1999415968 Cathepsin Y (a novel thiol enzyme) produces kinin potentiating peptide from the component protein of rat plasma Sakamoto E.; Sakao Y.; Taniguchi Y.; Yamafuji K. E. Sakamoto, Department of Food and Nutrition, Nakamura Gakuen University, Jonan-ku, Fukuoka 814-0198 Japan Immunopharmacology (IMMUNOPHARMACOLOGY) (Netherlands)
CODEN: IMMUD ISSN: 0162-3109
Publisher Item Identifier: S016231099900079X 1999 , 45/1-3 (207-214) Document Type: Journal; Article Summary Language: ENGLISH Language: ENGLISH Number of References: 18 Cathepsin Y (a novel thiol enzyme) produces kinin potentiating peptide from the component protein of rat plasma

Page 28

Rat spleen cathepsin Y (a novel enzyme) that produces bradykinin (BK) potentiating

peptide (BPP) from rat plasma was isolated.....from cDNA cloned by reverse

cathepsinsearch.txt transcription-polymerase chain reaction (RT-PCR). We propose the name cathepsin Y for this enzyme considering its origin, characteristics and the amino acid sequence. BPP potentiates not.....when the level is doubled. The precursor proteins that produce BPP by the action of cathepsin Y are eluted into two fractions when the heated plasma was applied to a negative ion....this paper, we report on the characteristics and the amino acid sequence of rat spleen cathepsin Y, its structure and the potentiating activity of BPP, and isolation of the precursor protein. Copyright.. DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *bradykinin--endogenous compound--ec; *thiolproteinase--endogenous compound--ec; * kinin--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin y--endogenous compound--ec 4/3,K/51 (Item 50 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1999379729 Cathepsin P, a novel protease in mouse placenta Sol-Church K.; Frenck J.; Troeber D.; Mason R.W. R.W. Mason, Laboratory of Enzymology, Department of Research, Alfred I. duPont Hospital Children, PO Box 269, Wilmington, DE 19899 United States Author Email: rmason@nemours.org Biochemical Journal (BIOCHEM. J.) (United Kingdom) 15 OCT 1999 , 343/2 (307 - 309)CODEN: BIJOA ISSN: 0264-6021 Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 13 Cathepsin P, a novel protease in mouse placenta The complete cDNA nucleotide sequence of a novel cathepsin derived from mouse placenta, termed cathepsin P, was determined. mRNA for cathepsin P was expressed in placenta and at lower levels in visceral yolk sac, but could not... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec proteinase--endogenous compound--ec; complementary DNA--endogenous compound--ec; messenger RNA--endogenous compound--ec 4/3,K/52 (Item 51 from file: 73) Links Fulltext available through: USPTO Fu USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1999344211 Human cathepsin X: A cysteine protease with unique carboxypeptidase activity Nagler D.K.; Zhang R.; Tam W.; Sulea T.; Purisima E.O.; Menard R. R. Menard, Biotechnology Research Institute, National Research Council of Canada, 6100 Royalmount Ave., Montreal, Que. H4P 2R2 Canada
Author Email: robert.menard@nrc.ca
Biochemistry (BIOCHEMISTRY) (United States) 28 SEP 1999 , 38/39 (12648-12654)

Cathepsin X is a novel cysteine protease which was identified recently from the EST (expressed sequence tags) database. In a homology model of the mature cathepsin X, a unique three residue insertion between the Gln22 of the oxyanion hole and the Page 29

Human cathepsin X: A cysteine protease with unique carboxypeptidase activity

ISSN: 0006-2960

Summary Language: ENGLISH

Document Type: Journal; Article

CODEN: BICHA

Language: ENGLISH

Number of References: 40

cathepsinsearch.txt active... ...verify this hypothesis, human procathepsin X was expressed in Pichia pastoris and converted to mature cathepsin X using small amounts of human cathepsin L. Cathepsin X was found to display excellent carboxypeptidase activity against the substrate Abz-FRF(4NOinf 2), with.....1 ssup -sup 1 at the optimal pH of 5.0. However, the activity of cathepsin X against the substrate Cbz-FR-MCA and Abz-AFRSAAQ-EDDnp was found to be extremely....k(cat)/K(M) values lower than 70 Msup -sup 1 ssup -sup 1. Therefore, cathepsin X displays a stricter exopeptidase activity than cathepsin B. No inhibition of cathepsin X by cystatin C could be detected up to a concentration of 4 muM of inhibitor....the bound carboxypeptidase substrate is predicted to establish a number of favorable contacts within the cathepsin X binding site, in particular with residues His23 and Tyr27 from the mini-loop. The presencesubstrates in the primed subsites of the protease. The marked structural and functional differences of cathepsin X relative to other members of the papain family of cysteine proteases will be of great... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec; *carboxypeptidase--endogenous compound--ec; * cathepsin l--endogenous compound--ec; *cathepsin b--endogenous compound--ec histidine--endogenous compound--ec; tyrosine--endogenous compound--ec; cystatin

c--endogenous compound--ec; cysteine Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/53 (Item 52 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1998297006 07403528 Human cathepsin X: A novel cysteine protease of the papain family with a very short proregion and unique insertions

Nagler D.K.; Menard R. R. Menard, Biotechnology Research Institute, National Research Council Canada, 6100 Avenue Royalmount, Montreal, Que. H4P 2R2 FEBS Letters (FEBS LETT.) (Netherlands) CODEN: FEBLA ISSN: 0014-5793 Canada 1998 , 434/1-2 (135-139) Publisher Item Identifier: S0014579398009648 Document Type: Journal ; Article Summary Language: ENGLISH Language: ENGLISH Number Of References: 40

Human cathepsin X: A novel cysteine protease of the papain family with a very short proregion and unique...

A novel cDNA encoding a cysteine protease of the papain family named cathepsin X was obtained by PCR amplification from a human ovary CDNA library. The cathepsin X cDNA is ubiquitously expressed in human tissues and contains an open reading frame of 912....highly conserved regions in papain-like cysteine proteases including the catalytic residues are present in cathepsin X. The mature part of cathepsin X is 26-32% identical to human cathepsins B, C, H, K, L, O, S and W. The cathepsin X sequence contains several unique features: (i) a very short proregion; (ii) a three amino acid...

DRUG DESCRIPTORS: * cathepsin: *papain--endogenous compound--ec; *cysteine proteinase --endogenous compound--ec cathepsin s--endogenous compound--ec; cathepsin b--endogenous compound--ec

4/3,K/54 (Item 53 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1998227542 07356252

Cathepsin Z, a novel human cysteine proteinase with a short propeptide domain and a Page 30

unique chromosomal location

Santamaria I.; Velasco G.; Pendas A.M.; Fueyo A.; Lopez-Otin C.
C. Lopez-Otin, Depto. de Bioquimica/Biologia Molec., Facultad de Medicina,
Universidad de Oviedo, 33006 Oviedo Spain
Author Email: CLO@DWARF1.QUIMICA.UNIOVI.ES
Journal of Biological Chemistry (J. BIOL. CHEM.) (United States) 03 JUL 1998 ,
273/27 (16816-16823)
CODEN: JBCHA ISSN: 0021-9258
Document Type: Journal; Article
Language: ENGLISH Summary Language: ENGLISH
Number Of References: 62
Cathepsin Z, a novel human cysteine proteinase with a short propeptide domain and a unique chromosomal location

...revealed that the isolated cDNA codes for a polypeptide of 303 amino acids, tentatively called cathepsin Z, that exhibits structural features characteristic of cysteine proteinases. Fluorescent in situ hybridization experiments revealed that the human cathepsin Z gene maps to chromosome 20q13, a location that differs from all cysteine proteinase genes mapped to date. The cDNA encoding cathepsin Z was expressed in Escherichia coli as a fusion protein with glutathione S- transferase, and after... ...amido-4- methylcoumarin, used as a substrate for cysteine proteinases. Northern blot analysis demonstrated that cathepsin Z is widely expressed in human tissues, suggesting that this enzyme could be involved in the normal intracellular protein degradation taking place in all cell types. Cathepsin Z is also ubiquitously distributed in cancer cell lines and in primary tumors from different sources... ...unusual short propeptide, together with its unique chromosomal location among cysteine proteinases, we propose that cathepsin Z may be the first representative of a novel subfamily of this class of proteolytic enzymes. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin z--endogenous compound--ec

4/3,K/55 (Item 54 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
06200066 EMBASE No: 1995231005
Cloning and complete coding sequence of a novel human cathepsin expressed in giant cells of osteoclastomas

Li Y.-P.; Alexander M.; Wucherpfennig A.L.; Yelick P.; Chen W.; Stashenko P. Forsyth Dental Center, 140 Fenway, Boston, MA 02115 United States Journal of Bone and Mineral Research (J. BONE MINER. RES.) (United States) 1995 , 10/8 (1197-1202) CODEN: JBMRE ISSN: 0884-0431 Document Type: Journal ; Article

Language: ENGLISH Summary Language: ENGLISH

...has been identified by differential screening of a human osteoclastoma cDNA library. This molecule, termed cathepsin X, appears to represent the human homolog of the osteoclast-expressed rabbit cathepsin OC-2. Cathepsin X (GenBank accession number U20280) is 93.9% identical to OC-2 at the amino acid level, and is 92% identical at the nucleotide level within the coding region. Cathepsin X is 52.2 and 46.9% identical to cathepsins S and L, respectively, and is therefore clearly distinct from these enzymes. Cathepsin X mRNA was localized to multinucleated giant cells within the osteoclastoma tumor by in situ hybridization. These data strongly support the hypothesis that cathepsin X represents a novel cysteine proteinase which is expressed at high levels in osteoclasts. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec cysteine proteinase--endogenous compound--ec; unclassified drug Page 31

Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/56 (Item 1 from file: 35) Links

Dissertation Abs Online

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01980017 ORDER NO: AADAA-IMQ83852

The design of substrates for cathepsin X

Author: Devanathan, Gopal

Degree: M.Sc. Year: 2003

Corporate Source/Institution: Concordia University (Canada) (0228)

Source: Volume 42/03 of MASTERS ABSTRACTS. of Dissertations Abstracts International.

PAGE 941 . 90 PAGES ISBN: 0-612-83852-8

The design of substrates for cathepsin X

...diseases such as arthritis, Alzheimer's, and cancer, they are attractive targets for inhibitor design. Cathepsin X is a cysteine protease that was only recently discovered. The primary structure of cathepsin X contains several unique features that clearly distinguish it from the other human cysteine proteases. The... ... a systematic study on the S2, S1, and S1<super>′</super> subsites of the cathepsin X active site and to gain a detailed understanding of the enzyme's substrate specificity.

Three libraries of compounds have been synthesized based on the parent compound 2-Abz-Phe-Arg-Phe(4NO₂). In each library, the 20 natural... ...prime;</super> sites respectively, while keeping the other positions fixed. In reference to the parent compound, P2 is occupied by Phe, P1 by Arg, and P1<super>′</super> by Phe.....by docking 2-Abz-Phe-Arg-Phe(4NO₂) and analogues to the cathepsin X active site in order to gain a detailed understanding of factors underlying substrate specificity. Knowledge...

4/3,K/57 (Item 1 from file: 357) Links
Derwent Biotech Res.
(c) 2007 The Thomson Corp. All rights reserved.
0377298 DBA Accession No.: 2005-23004 PATENT
Identification of a compound capable of modulating the activity

Identification of a compound capable of modulating the activity of cathepsin Z in a cell comprises measuring the cell's base level of cathepsin Z activity in the absence and presence of the compound involving vector-mediated gene transfer and expression in host cell for therapy

Author: SALTZMAN A G; TANG Z; PALEJWALA V; CAVALLO J

Patent Assignee: AVENTIS PHARM INC 2005

Patent Number: WO 200565693 Patent Date: 20050721 WPI Accession No.: 2005-533570 (200554)

Priority Application Number: US 533330 Application Date: 20031230 National Application Number: WO 2004US41815 Application Date: 20041214

Language: English
Identification of a compound capable of modulating the activity of cathepsin Z in a cell comprises measuring the cell's base level of cathepsin Z activity in the absence and presence of the compound involving vector-mediated gene transfer and expression in host cell for therapy
Abstract: DERWENT ABSTRACT: NOVELTY - Identification of a compound (I) capable of

Abstract: DERWENT ABSTRACT: NOVELTY - Identification of a compound (I) capable of modulating the activity of cathepsin Z in a cell comprises measuring the cell's base level of cathepsin Z activity in the absence of a candidate compound; introducing the candidate compound; and measuring the cell's level of cathepsin Z activity in the presence of the candidate compound. DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for: (1) the compound (I) capable of modulating activity of cathepsin Z; and (2) a pharmaceutical comprising (I) and excipient for treating an inflammatory. ACTIVITY - Antiinflammatory; Immunosuppressive; Antiarthritic;

Page 32

Antirheumatic: Neuroprotective. MECHANISM OF ACTION - Cathepsin Z modulator. Test details are described but no results given. USE - (I) is useful to treat...

Descriptors: recombinant cathepsin-Z prep., isol., vector-mediated gene transfer, expression in host cell, appl., inflammatory disease, autoimmune disease...

4/3,K/58 (Item 2 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0338230 DBA Accession No.: 2004-10522 PATENT Use of polynucleotide sequence encoding Cathespin Y protein for identification of therapeutic agent useful for treating stroke e.g. ischemic stroke vector-mediated cathespin-Y gene transfer, expression in host cell and antisense oligonucleotide for drug screening and gene therapy

Author: LUBBERT H; ZWILLING S; ENGELS P

Patent Assignee: LUBBERT H; ZWILLING S; ENGELS P 2003 Patent Number: US 20030232740 Patent Date: 20031218 WPI Accession No.: 2004-142033 (200414)

Priority Application Number: US 392809 Application Date: 20030319 National Application Number: US 392809 Application Date: 20030319

Language: English

Abstract: ...of potential therapeutic agent for treating stroke involves contacting a cell capable of expressing a Cathepsin Y gene or homologues or fragments with the potential therapeutic agent; detecting a level of expression of the Cathepsin Y gene in the test cell; comparing expression in the test cell to a reference cell... potential therapeutic agent for treating stroke involves contacting a cell capable of expressing a Cathepsin Y gene or homologues or fragments with the potential therapeutic agent; detecting a level of expression of the Cathepsin Y gene in the test cell; comparing the level of expression of the Cathepsin Y gene in the test cell to a level of expression of the Cathepsin Y gene in the test cell to a level of expression of the Cathepsin Y gene in a reference cell whose disease stage is known; and identifying the difference in the expression level of the Cathepsin Y gene in the test cell and the reference cell. INDEPENDENT CLAIMS are included for the following: (a) a composition comprising a compound of formula (I) or its salt; (b) a composition comprising a nucleic acid sequence (S1) which is an anticome compared to a nucleic acid sequence (S2) encoding Cathepsin Y antisense sequence compared to a nucleic acid sequence (S2) encoding Cathepsin Y, its homologue or fragment. (S2) Has sequence of 1140 or 1500 nucleotide bases as given... is 0, then R4 is other than -N(CH3)OCH3. ACTIVITY - Cerebroprotective. MECHANISM OF ACTION - Cathepsin Y protein inhibitor. Test details are described, but no results are given. USE - For identifying potential...

4/3,K/59 (Item 3 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0335363 DBA Accession No.: 2004-07655 PATENT New antisense compound targeted to nucleic acid molecules encoding cathepsin Z, useful for treating diseases associated with expression of cathepsin Z, e.g. encephalitis, viral infection, or hyperproliferative disorder involving vector-mediated gene transfer and expression in host cell for use in therapy

Author: DOBIE K W

Patent Assignee: ISIS PHARM INC 2003

Patent Number: US 20030224511 Patent Date: 20031204 WPI Accession No.: 2004-060543

Priority Application Number: US 159266 Application Date: 20020531 National Application Number: US 159266 Application Date: 20020531

Language: English

New antisense compound targeted to nucleic acid molecules encoding cathepsin Z, useful for treating diseases associated with expression of cathepsin Z, e.g. encephalitis, viral infection, or hyperproliferative disorder involving vector-mediated gene transfer and expression...

Abstract: DERWENT ABSTRACT: NOVELTY - A company (I) 8-80 nucleobases in length

Page 33

targeted to a nucleic acid molecule encoding cathepsin Z, is new. DETAILED DESCRIPTION - A compound (I) 8-80 nucleobases in length targeted to a nucleic acid molecule encoding cathepsin Z, is new. The compound specifically hybridizes with the nucleic acid molecule encoding cathepsin Z and inhibits the expression of cathepsin Z, or specifically hybridizes with at least an 8-nucleobase portion of a preferred target region on a nucleic acid molecule encoding cathepsin Z. INDEPENDENT CLAIMS are included for the following: (1) a composition comprising (I) and a pharmaceutical carrier or diluent; (2) a method of inhibiting the expression of cathepsin Z in cells or tissues comprising contacting the cells or tissues with (I); and (3) a method of treating an animal having a disease or condition associated with cathepsin Z comprising administering to the animal a therapeutic or prophylactic amount of (I) so that expression of cathepsin Z is inhibited. BIOTECHNOLOGY - Preparation: The antisense compounds are produced by solid phase synthesis. Preferred Compound: The compound is an antisense oligonucleotide, preferably a chimeric oligonucleotide. The antisense oligonucleotide comprises: (a) at least... ...Inhibitor Z. USE - The antisense oligonucleotides and compounds are useful for inhibiting the expression of cathepsin Z, and for treating diseases or conditions associated with expression of cathepsin Z, e.g. encephalitis, viral infection, or hyperproliferative disorder, such as cancer (all claimed). The antisense... Descriptors: recombinant cathepsin-Z prep., isol., vector-mediated gene transfer, expression in host cell, antisense oligonucleotide, appl. encephalitis, virus...

4/3,K/60 (Item 4 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0334375 DBA Accession No.: 2004-06667 PATENT Composition useful for treating pain e.g. neuropathic pain comprises polynucleotide sequence sense and antisense sequence for use in disease therapy and gene therapy

Author: LUBBERT H; ENGELS P; SCHMITZ B Patent Assignee: LUBBERT H; ENGELS P; SCHMITZ B 2003
Patent Number: US 20030212003 Patent Date: 20031113 WPI Accession No.: 2004-041675

Priority Application Number: US 369386 Application Date: 20030214 National Application Number: US 369386 Application Date: 20030214

Language: English

Abstract: ...therapeutic agents for treating pain involving: (a) contacting a test cell capable of expressing a Cathepsin Y gene, its homologues or fragments with the potential therapeutic agent; (b) detecting a level of expression of the Cathepsin Y gene in the test cell; (c) comparing the level of expression of the Cathepsin Y gene in the test cell to that in a reference cell; and (d) identifying the difference in the expression levels of the Cathepsin Y gene in the test cell and reference cell; (2) identification of a therapeutic agent for treating pain involving: (a) incubating a sample comprising a Cathepsin Y protein, a test compound/agent and a polypeptide which is a target of Cathepsin Y protein proteolysis; (b) determining an aminotorminal amino acid of a portion resulting from the proteolysis. aminoterminal amino acid of a peptide resulting from the proteolysis... ...nucleic acid sequence is an antisense sequence compared to a nucleic acid sequence that encodes Cathepsin Y and has a sequence of 1387 or 1500 nucleotide bases. BIOTECHNOLOGY - Preferred Method: The expression of the Cathepsin Y gene is determined by at least one method selected from PCR of a cDNA, hybridizing a sample DNA and detecting a Cathepsin Y protein. ACTIVITY - Analgesic; Antidiabetic; Neuroprotective; Virucide; Vulnerary; Cardiant. MECHANISM OF ACTION - Cathepsin Y protein inhibitor. No biological data given. USE - The compound is useful for treating pain e.g. neuropathic pain (claimed), diabetic neuropathy, post-herpetic neuralgia.....reflex sympathetic dystrophy and causalgia, myocardial syndromes or idionathic pain. ADVANTAGE - The composition efficiently downregulates Cathensin Y idiopathic pain. ADVANTAGE - The composition efficiently downregulates Cathepsin Y activity and hence treats pain. EXAMPLE - No relevant example given. (22 pages) Descriptors: polynucleotide composition, cathepsin Y gene, antisense sequence, polymerase chain reaction, appl. pain, neuropathic pain, diabetic neuropathy, post-herpetic neuralgia...

4/3,K/61 (Item 5 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0328457 DBA Accession No.: 2004-00749 PATENT Developing medicament used for treating pain comprises using polynucleotide sequence encoding cathepsin Y involving vector-mediated gene transfer and expression in host cell for use in neuropathic pain therapy Author: LUEBBERT H; SCHMITZ B Patent Assignee: BIOFRONTERA PHARM AG 2003 Patent Number: EP 1336847 Patent Date: 20030820 WPI Accession No.: 2003-814978 (2003) Priority Application Number: EP 20023400 Application Date: 20020214 National Application Number: EP 20023400 Application Date: 20020214 Language: English Developing medicament used for treating pain comprises using polynucleotide sequence encoding cathepsin Y involving vector-mediated gene transfer and expression in host cell for use in neuropathic pain... Abstract: DERWENT ABSTRACT: NOVELTY - Developing a medicament for treating pain comprises using a polynucleotide sequence encoding cathepsin Y. DETAILED DESCRIPTION - Developing a medicament for treating pain, for diagnosing pain status outside of a....efficacy of pain treatment outside of a living body, comprises using a polynucleotide sequence encoding cathepsin Y or homologs or fragments or the corresponding protein or peptide. INDEPENDENT CLAIMS are also included for: (1) the use of a compound downregulating cathepsin Y expression or activity for manufacture of a medicament for treatment of pain; (2) a diagnostic....isolated nucleic acid sequence comprising an 'antisense' sequence compared to a nucleic acid sequence encoding cathepsin Y or a fragment of the nucleic acid sequence as a medicament; and (4) a transgenic animal where the gene encoding cathepsin Y is manipulated in the animal in comparison to the wild type. ACTIVITY - Analgesic. MECHANISM OF ACTION -Cathepsin Y inhibitor. Tests are described, but no results are given. USE - Used for treating pain, particularly... Descriptors: recombinant cathepsin-Y prep., isol., vector-mediated gene transfer, expression in host cell, polymerase chain reaction, appl. neuropathic... 4/3,K/62 (Item 6 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0313249 DBA Accession No.: 2003-14389 PATENT New transgenic mice comprising a disruption in a cathepsin Z (CTSZ) useful as models for diseases or conditions associated with phenotypes relating to a disruption in a CTSZ gene, and in identifying drugs for treating a disease vector-mediated mutant gene transfer and expression in embryonic stem cell for transgenic mouse construction for use as an animal model in disease therapy Author: WISOTZKEY R G; KIRK C J Patent Assignee: DELTAGEN INC 2003

Patent Number: WO 200326403 Patent Date: 20030403 WPI Accession No.: 2003-354621

(200333) Priority Application Number: US 324639 Application Date: 20010924 National Application Number: WO 2002US30506 Application Date: 20020924 Language: English

New transgenic mice comprising a disruption in a cathepsin Z (CTSZ) useful as models for diseases or conditions associated with phenotypes relating to a disruption... Abstract: DERWENT ABSTRACT: NOVELTY - A transgenic mouse comprising a disruption in a cathepsin Z (CTSZ) gene, where there is no native expression of endogenous CTSZ gene, is new. DETAILED... ... a pharmaceutical composition for a condition associated with a function of CTSZ, comprises identifying a compound that modulates CTSZ, synthesizing the identified compound, and incorporating the compound into a pharmaceutical carrier. USE - The transgenic mouse is useful as a model for diseases....symptoms; and in testing and developing new treatments relating to behavioral phenotypes. EXAMPLE - Disruptions in cathepsin Z (CTSZ) genes were

produced by homologous recombination. Transgenic mice comprising disruptions in CTSZ genes were.. Descriptors: transgenic mouse construction, vector-mediated mutant cathepsin- Z gene transfer, expression in embryonic stem cell, phenotyping, animal model, antagonist, agonist, database, homologous recombination... 4/3,K/63 (Item 7 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0257611 DBA Accession No.: 2000-12101 PATENT New human cathepsin-Y protein, a gene encoding it and its application - diagnosis, therapy, gene therapy and drug screening Corporate Source: Japan. 2000 Patent Assignee: Fuii-Pharm. Patent Number: JP 2000157263 Patent Date: 20000613 WPI Accession No.: 2000-468198 (2041)Priority Application Number: JP 98352110 Application Date: 19981126 National Application Number: JP 98352110 Application Date: 19981126 Language: Japanese New human cathepsin-Y protein, a gene encoding it and its application Abstract: A human-derived cathepsin-Y protein (I) or a new human-derived cathepsin-Y protein which has at least 49% homology to the protein sequence of (I) has a...
...I) or its salts, peptides, etc., the DNA or the antibody; a drug containing a compound promoting or inhibiting biological activity of one of the claimed proteins, their partial peptides or... Descriptors: human recombinant cathepsin-Y prep., cysteine protease act., monoclonal antibody, vector expression in host cell, DNA probe hybridization, appl... ? s cathepsin **S1** 88555 S CATHEPSIN s cathepsin(w)Z or CSTZ or cathepsin(w)X or cathepsin(w)P or cathepsin(w)Y Processing Processing Processing 88555 CATHEPSIN 1099189 152 CATHEPSIN(W)Z 10 CSTZ 88555 **CATHEPSIN** 6940523 CATHEPSIN(W)X 197 88555 CATHEPSIN 12346779 .65 CATHEPSIN(W)P 88555 **CATHEPSIN** 1989729 CATHEPSIN(W)Y 30 S2 450 S CATHEPSIN(W)Z OR CSTZ OR CATHEPSIN(W)X OR CATHEPSIN(W)P OR CATHEPSIN(W)Y s s2 and compound 450 S2 4641049 **COMPOUND** S S2 AND COMPOUND **S**3 67 ? rd

Duplicate detection is not supported for File 393.

Page 36

>>>W:

cathepsinsearch.txt Duplicate detection is not supported for File 391. Records from unsupported files will be retained in the RD set. 63 RD (UNIQUE ITEMS) ? t s64/3, k/1-63>>>E: Set 64 does not exist ? t s4/3, k/1-63>>>w: KWIC option is not available in file(s): 399 4/3,K/1 (Item 1 from file: 5) Links USPTO Full Text Retrieval Options Fulltext available through: Biosis Previews(R) (c) 2007 The Thomson Corporation. All rights reserved. Biosis No.: 200510043834 18349334 Defining the substrate specificity of mouse cathepsin P Author: Puzer Luciano; Barros Nilana M T; Oliveira Vitor; Julianoa Maria Aparecida; Lu Guizhen: Hassanein Mohamed; Juliano Luiz; Mason Robert W; Carmona Adriana K (Reprint) Author Address: UNIFESP, Escola Paulista Med, Dept Biophys, Rua Tres Maio 100, BR-04044020 Sao Paulo, Brazil**Brazil Author E-mail Address: adriana@biofis.epm.br Journal: Archives of Biochemistry and Biophysics 435 (1): p 190-196 MAR 1 05 2005 ISSN: 0003-9861 Document Type: Article Record Type: Abstract Language: English Defining the substrate specificity of mouse cathepsin P Abstract: Cathepsin P is a recently discovered placental cysteine protease that is structurally related to the more ubiquitously expressed, broad-specificity enzyme, cathepsin L. We studied the substrate specificity requirements of recombinant meuse cathepsin P using fluorescence resonance energy transfer (FRET) peptides derived from the lead sequence Abz-KLRSSKQ-EDDnp......Arg), and hydrophobic aliphatic or aromatic residues (Val, Phe). For several substrates, the activity of cathepsin P was markedly regulated by kosmotropic salts, particularly Na2SO4. No significant effect on secondary or tertiary.....this substrate was almost two orders of magnitude higher than that of the original parent compound. These results show that cathepsin P, in contrast to other mammalian cathepsins, has a restricted catalytic specificity. (C) 2004 Elsevier Inc... Registry Numbers: ...cathepsin P **DESCRIPTORS:** Chemicals & Biochemicals: ...cathepsin P--4/3,K/2 (Item 1 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. 14746499 EMBASE No: 2007432049 Differential Expression of Cathepsins and Cystatin C in Ovine Uteroplacental Tissues Song G.; Bazer F.W.; Spencer T.E. T.E. Spencer, Center for Animal Biotechnology and Genomics, Department of Animal Science, Texas A and M University, 2471 TAMU, College Station, TX 77843-2471 **United States** Author Email: tspencer@tamu.edu Placenta (PLACENTA) (United Kingdom) 2007 , 28/10 (1091-1098) CODEN: PLACD ISSN: 0143-4004 Publisher Item Identifier: S0143400407001099 Document Type: Journal ; Article

Page 37

```
Language: ENGLISH Summ
Number Of References: 34
                           Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cystatin C--endogenous compound--ec
cathepsin B--endogenous compound--ec; cathepsin D--endogenous compound--ec;
cathepsin H--endogenous compound--ec; cathepsin K--endogenous compound--ec;
cathepsin L--endogenous compound --ec; cathepsin S--endogenous compound--ec;
messenger RNA --endogenous compound--ec; peptide hydrolase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec
 4/3,K/3 (Item 2 from file: 73)
                                             Links
    Fulltext available through:
                                             USPTO Full Text Retrieval Options
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                  EMBASE No: 2007348195
14643786
Expression of Cathepsin P mRNA, Protein and Activity in the Rat Choriocarcinoma Cell
Line, Rcho-1, During Giant Cell Transformation
Hassanein M.; Korant B.D.; Lu G.; Mason R.W.
R.W. Mason, Department of Biomedical Research, Alfred I duPont Hospital for
Children, 1600 Rock land Road, Wilmington, DE 19803 United States
Author Email: rmason@nemours.org
Placenta ( PLACENTA ) ( United Kingdom ) CODEN: PLACD ISSN: 0143-4004
                                                           2007 , 28/8-9 (912-919)
Publisher Item Identifier: S0143400406002773
Document Type: Journal; Article
Language: ENGLISH
                           Summary Language: ENGLISH
Number Of References: 44
Expression of Cathepsin P mRNA, Protein and Activity in the Rat Choriocarcinoma Cell
Line, Rcho-1, During Giant Cell...
...proteases perform critical functions in protein turnover and are essential for normal growth and development. Cathepsin P is a member of a newly discovered family of lysosomal cysteine proteases uniquely expressed in... L was not regulated. A specific enzyme assay was developed to show that activity of cathepsin P mirrored mRNA expression during differentiation. Cathepsin P protein co-localizes with
cathepsin B, indicating that the enzyme probably functions in the endosomal ...
DRUG DESCRIPTORS:
  cathepsin--endogenous compound--ec
cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec; peptide
hydrolase--endogenous compound--ec; proteinase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin 1--endogenous compound--ec; cathepsin
2--endogenous compound--ec; cathepsin P--endogenous compound --ec; cathepsin
Q--endogenous compound--ec; cathepsin m--endogenous compound--ec; cathepsin
r--endogenous compound--ec
 4/3,K/4 (Item 3 from file: 73)
                                             Links
    Fulltext available through:
                                             USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved. 14625291 EMBASE No: 2007394692
Inflammatory processes in the aging mouse brain: Participation of dendritic cells
and T-cells
Stichel C.C.; Luebbert H.
C.C. Stichel, Biofrontera Bioscience GmbH, D-51377 Leverkusen
                                                                                        Germany
Author Email: c.stichel-gunkel@biofrontera.com
Neurobiology of Aging ( NEUROBIOL. AGING ) ( United States )
                                                                                      2007 , 28/10
(1507-1521)
CODEN: NEAGD
                    ISSN: 0197-4580
Publisher Item Identifier: S0197458006002740
                                                     Page 38
```

```
Document Type: Journal ; Article
Language: ENGLISH Summa
Number Of References: 100
                        Summary Language: ENGLISH
DRUG DESCRIPTORS:
CD11b antigen--endogenous compound--ec; cathepsin S--endogenous compound--ec;
cathepsin--endogenous compound--ec; integrin --endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/5 (Item 4 from file: 73)
                                        Links
    Fulltext available through:
                                        USPTO Full Text Retrieval Options
EMBASE
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14423842 EMBASE No: 2007158978
Differential expression of cathepsin X in aging and pathological central nervous
system of mice
Wendt W.; Zhu X.-R.; Lubbert H.; Stichel C.C.
C.C. Stichel, Biofrontera Bioscience GmbH, D-51377 Leverkusen
                                                                               Germany
Author Email: c.stichel-qunkel@biofrontera.com
Experimental Neurology (EXP. NEUROL.) (United States)
CODEN: EXNEA ISSN: 0014-4886 eISSN: 1090-2430
Publisher Item Identifier: S0014488607000222
                                                                          2007 , 204/2 (525-540)
Document Type: Journal ; Article
Language: ENGLISH Summ
Number Of References: 77
                        Summary Language: ENGLISH
Differential expression of cathepsin X in aging and pathological central nervous
system of mice
...we analyzed the regional, cellular and subcellular localization and the activity
of the recently discovered cathepsin X in the normal, developing and pathological
mouse brain. Our results show that CATX is: (i... ...plaques in a transgenic mouse model and in Alzheimer patients. These results strongly suggest that cathepsin X is an important player in degenerative processes during normal aging and in pathological conditions. (c...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/6 (Item 5 from file: 73)
                                        Links
   Fulltext available through:
                                        USPTO Full Text Retrieval Options
EMBASE
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                EMBASE No: 2006599352
14198060
Cysteine cathepsins: Regulators of antitumour immune response
Obermajer N.; Doljak B.; Kos J.
J. Kos, University of Ljubljana, Department of Pharmaceutical Biology, Faculty of
Pharmacy, Askerceva 7, SI-1000 Ljubljana
                                                    Slovenia
Author Email: Janko.kos@ffa.uni-lj.si
Expert Opinion on Biological Therapy ( EXPERT OPIN. BIOL. THER. ) ( United Kingdom )
  2006 , 6/12 (1295-1309)
CODEN: EOBTA
                  ISSN: 1471-2598
Document Type: Journal ; Review
Language: ENGLISH Summar
Number Of References: 120
                        Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
cysteine derivative--endogenous compound--ec; cysteine proteinase --endogenous
compound--ec; major histocompatibility antigen class 2 --endogenous compound--ec;
cytokine--endogenous compound--ec; growth factor--endogenous compound--ec; integrin--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin
                                               Page 39
```

```
cathepsinsearch.txt
L--endogenous compound--ec; cathepsin S--endogenous compound --ec; cathepsin
K--endogenous compound--ec; stefin A--endogenous compound--ec; stefin B--endogenous
compound--ec; cystatin C --endogenous compound--ec; cystatin--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin H--endogenous
Drug Terms (Uncontrolled): cathepsin w--endogenous compound--ec; cathepsin x
--endogenous compound--ec; cystatin f--endogenous compound --ec; cathepsin
o--endogenous compound--ec; cathepsin v--endogenous compound--ec
 4/3,K/7 (Item 6 from file: 73)
    Fulltext available through:
                                       USPTO Full Text Retrieval Options
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14090484 EMBASE No: 2006493081
Cargo selectivity of the ERGIC-53/MCFD2 transport receptor complex
Nyfeler B.; Zhang B.; Ginsburg D.; Kaufman R.J.; Hauri H.-P.
H. Hans-Peter, Biozentrum, University of Basel, CH-4056 Basel Author Email: hans-peter.hauri@unibas.ch
                                                                            Switzerland
Traffic (TRAFFIC) (Denmark) 2006, 7/11 (14 CODEN: TRAFF ISSN: 1398-9219 eISSN: 1600-0854
                                                7/11 (1473-1481)
CODEN: TRAFF ISSN: 1398-9219 (
Document Type: Journal; Article
Language: ENGLISH
                      Summary Language: ENGLISH
Number Of References: 37
...complex in the early secretory pathway. ERGIC-53 also interacts with the two lysosomal glycoproteins cathepsin Z and cathepsin C. Here, we tested the subunit
interdependence and cargo selectivity of ERGIC-53... ... yellow fluorescent protein
fragment complementation. We found that MCFD2 is dispensable for the binding of
cathepsin Z and cathepsin C to ERGIC-53. The results indicate that ERGIC-53 can bind
DRUG DESCRIPTORS:
* endoplasmic reticulum golgi intermediate compartment protein 53--endogenous
compound--ec; *protein--endogenous compound--ec
secretory protein--endogenous compound--ec; receptor--endogenous compound--ec; blood
clotting factor 5--endogenous compound --ec; blood clotting factor 8--endogenous
compound--ec; lectin --endogenous compound--ec; glycoprotein; dipeptidyl peptidase I; cathepsin; protein subunit--endogenous compound--ec; small interfering RNA;
yellow fluorescent protein; unclassified drug
Drug Terms (Uncontrolled): multiple coagulation factor deficiency protein
2--endogenous compound--ec
 4/3,K/8 (Item 7 from file: 73)
                                      Links
   Fulltext available through:
                                       USPTO Full Text Retrieval Options
EMBASE
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               EMBASE No: 2006471802
14057877
Cysteine cathepsins: Multifunctional enzymes in cancer
Mohamed M.M.; Sloane B.F.
B.F. Sloane, Department of Pharmacology, Wayne State University School of Medicine,
Detroit, MI 48201 United States
Author Email: bsloane@med.wayne.edu
                                                                           2006 , 6/10
Nature Reviews Cancer ( NAT. REV. CANCER ) ( United Kingdom )
(764 - 775)
                ISSN: 1474-175X
CODEN: NRCAC
Publisher Item Identifier: NRC1949
Document Type: Journal; Conference Paper
Language: ENGLISH Summa
Number Of References: 145
                      Summary Language: ENGLISH
```

DRUG DESCRIPTORS:

```
cathepsinsearch.txt
* cysteine--endogenous compound--ec; *cathepsin--endogenous compound--ec cathepsin B--endogenous compound--ec; dipeptidyl peptidase I --endogenous
compound--ec; cathepsin H--endogenous compound --ec; cathepsin L--endogenous
compound--ec; cathepsin K--endogenous compound--ec; cathepsin S--endogenous
compound--ec; kiningen --endogenous compound--ec; cystatin C--endogenous compound
--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin O--endogenous compound--ec; cathepsin
V--endogenous compound--ec; cathepsin W--endogenous compound--ec; cathepsin x--endogenous compound--ec; cystatin D --endogenous compound--ec; cystatin E--endogenous compound--ec; cystatin f--endogenous compound--ec; cystatin s--endogenous compound--ec; cystatin sn--endogenous compound--ec; cystatin sn--endogenous compound--ec; cystatin sn--endogenous compound--ec; cystatin sn--endogenous compound--ec
 4/3,K/9 (Item 8 from file: 73)
                                                Links
     Fulltext available through:
                                                USPTO Full Text Retrieval Options
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                   EMBASE No: 2006446396
14049123
Molecular aspects of stromal-parenchymal interactions in malignant neoplasms
A. Zalatnai, First Department of Pathology and Experimental Cancer Research, Semmelweis University, Faculty of Medicine, Ulloi ut 26, H-1085 Budapest H
                                                                                                             Hungary
Author Email: zalatnái@korb1.sote.hu
                                                                                               2006, 6/6
Current Molecular Medicine ( CURR. MOL. MED. ) ( Netherlands )
(685 - 693)
CODEN: CMMUB
                      ISSN: 1566-5240
Document Type: Journal; Review
Language: ENGLISH Summa
Number Of References: 108
                             Summary Language: ENGLISH
DRUG DESCRIPTORS:
cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec; enzyme
precursor--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin-endogenous compound--ec; matrix cathepsin-endogenous compound--ec; cathepsin-endogenous compound--ec; matrix cathepsin-endogenous compound--ec; cathepsin-endogenous compound--ec;
interleukin lalpha--endogenous compound--ec; basic fibroblast growth
factor--endogenous compound--ec; gelatinase B--endogenous compound--ec; tumor
necrosis factor alpha-endogenous compound-ec; transforming growth factor
beta--endogenous compound--ec; gelatinase A--endogenous compound--ec; collagenase
3--endogenous compound--ec; CD68 antigen--endogenous compound--ec; stromal cell
derived factor 1 --endogenous compound--ec; transforming growth factor beta1
--endogenous compound--ec; inducible nitric oxide synthase --endogenous
compound--ec; gemcitabine--pharmacology--pd; matrix metalloproteinase
inhibitor--clinical trial--ct; matrix metalloproteinase inhibitor--drug therapy...
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/10 (Item 9 from file: 73) Links
    Fulltext available through:
                                                USPTO Full Text Retrieval Options
EMBASE
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13924109 EMBASE No: 2006345822
Caenorhabditis elegans: Study model for animal and human cathepsins and inhibitors
Hashmi S.; Anwer K.; Bilgrami A.L.
S. Hashmi, Laboratory of Molecular Parasitology, Lindsley F. Kimbal Research
Instiute, New York Blood Center, 310 East 67th Street, New York, NY 10021
Author Email: shashmi@nybloodcenter.org
Current Enzyme Inhibition ( CURR. ENZYME INHIB. ) ( Netherlands ) 2006, 2/2
(173-188)
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Page 41

ISSN: 1573-4080 Document Type: Journal; Review Language: ENGLISH Summary Language: ENGLISH Number of References: 236 ...L enzymes in C. elegans. Besides, it also reviews the function of a recently described cathepsin Z. (c) 2006 Bentham Science Publishers Ltd. DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cysteine proteinase inhibitor --drug therapy--dt; *cysteine proteinase inhibitor--endogenous compound--ec; *cysteine proteinase inhibitor--pharmacology--pd; * cysteine proteinase inhibitor--topical drug administration--tp cathepsin L--endogenous compound--ec; cathepsin B--endogenous compound--ec; cysteine proteinase--endogenous compound--ec; papain--endogenous compound--ec; cathepsin E--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin H--endogenous compound --ec; cathepsin K--endogenous compound--ec; stefin A--endogenous compound--ec; stefin B--endogenous compound--ec; anthelmintic agent; cyclophosphamide--drug therapy--dt; cyclophosphamide--pharmacology --pd; antineoplastic agent--drug therapy--dt; antineoplastic agent --pharmacology--pd; proteinase inhibitor--drug therapy--dt; proteinase inhibitor--endogenous compound--ec; proteinase inhibitor--tp; antimalarial agent--drug therapy--dt; antimalarial agent--drug therapy--dt; antimalarial agent--pharmacology--pd; cystatin C--drug therapy--dt; cystatin C--endogenous compound--ec; cystatin C--pharmacology--pd; cystatin C--topical drug administration--tp; cystatin C--pharmacology--pd; cystatin C--topical drug administration--tp; cystatin--pharmacology--pd; antivirus agent--drug therapy--dt; antivirus oryzacystatin--pharmacology--pd; antivirus agent--drug therapy--dt; antivirus agent--endogenous compound--ec; antivirus agent --pharmacology--pd; antivirus agent--topical drug administration--tp; unindexed drug; unclassified drug Drug Terms (Uncontrolled): cathepsin inhibitor--drug therapy--dt; cathepsin inhibitor--endogenous compound--ec; cathepsin inhibitor--pharmacology--pd; cathepsin inhibitor--topical drug administration--tp; cathepsin Z --endogenous compound--ec; peptide aldehyde--pharmacology--pd; alpha ketoamide--drug therapy--dt... 4/3,K/11 (Item 10 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2006331563 13908004 Carboxypeptidase cathepsin X mediates betaSUB2-integrin-dependent adhesion of differentiated U-937 cells Obermajer N.; Premzl A.; Zavas(caron)nik Bergant T.; Turk B.; Kos J. J. Kos, Faculty of Pharmacy, University of Ljubljana, As(caron)kerc(caron)eva 7, SI-1000 Ljubljana Slovenia Author Email: janko.kos@ffa.uni-lj.si Experimental Cell Research (EXP. CELL RES.) (United States) 01 AUG 2006, 312/13 (2515-2527) CODEN: ECREA ISSN: 0014-4827 Publisher Item Identifier: S0014482706001601 Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 50 Carboxypeptidase cathepsin X mediates betaSUB2-integrin-dependent adhesion of differentiated U-937 cells Cathepsin X is a lysosomal carboxypeptidase with a potential role in processes of inflammation and immune response....integrin-binding motifs RGD and ECD, present in the pro- and in mature forms of cathepsin X, respectively, suggest that this enzyme might have a function in cell signaling and adhesion. In....protease inhibitors E-64 and CA-074 and 2F12 monoclonal antibody, all of which inhibit cathepsin X activity, significantly reduced adhesion of differentiated U-937 cells

Page 42

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cathepsinsearch.txt
to polystyrene- and fibrinogen-coated surfaces....whereas their binding to vitronectin, fibronectin or Matrigel was not affected. On the other hand, cathepsin X, added to differentiating U-937 cells, stimulated their adhesion. Using confocal microscopy, we demonstrated that the pro-form of cathepsin X was co-localized with
betaSUB2 and betaSUB3 integrin subunits and its mature form solely with... ... U-937
cells and in co-cultures with endothelial cells. Our results indicate that active
cathersin X mediates the function of betaSUB2 integrin receptors during cell
adhesion and that it could also...
DRUG DESCRIPTORS:
* carboxypeptidase--endogenous compound--ec; *beta2 integrin --endogenous
compound--ec
...cysteine proteinase inhibitor; monoclonal antibody; polystyrene; fibrinogen; integrin receptor; vitronectin; fibronectin; matrigel; beta3 integrin--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; ca 074
 4/3,K/12 (Item 11 from file: 73) Links
    Fulltext available through:
                                         USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                EMBASE No: 2006270160
Tumor cell-derived and macrophage-derived cathepsin B promotes progression and lung
metastasis of mammary cancer
Vasiljeva O.; Papazoglou A.; Kruger A.; Brodoefel H.; Korovin M.; Deussing J.;
Augustin N.; Nielsen B.S.; Almholt K.; Bogyo M.; Peters C.; Reinheckel T. T. Reinheckel, Institut fur Molekulare Medizin und Zellforschung,
Albert-Ludwigs-Universitat Freiburg, Stefan Meier Strasse 17, D-79104 Freiburg
Author Email: Thomas.Reinheckel@uniklinik-freiburg.de
Cancer Research ( CANCER RES. ) ( United States ) 15 MAY 2006 , 66/10 (5242-5250) CODEN: CNREA ISSN: 0008-5472
CODEN: CNREA 155N: 0000-3472
Document Type: Journal ; Article
Language: ENGLISH
                        Summary Language: ENGLISH
Number Of References: 48
...labeling of cysteine cathepsins by the active site probe DCG-04 detected
up-regulation of cathepsin X on PyMT;ctsbSUP+/+ cells. Treatment of cells with a
neutralizing anti-cathepsin X antibody significantly reduced Matrigel invasion of PyMT;ctsbSUP+/+ cells but did not affect invasion of PyMT;ctsbSUP+/+ or
PyMT; ctsbSUP+/- cells, indicating a compensatory function of cathepsin X in
CTSB-deficient tumor cells. Finally, an adoptive transfer model, in which ctsbSUP+/+, ctsbSUP+/-, and...
DRUG DESCRIPTORS:
* cathepsin B--endogenous compound--ec
virus middle T antigen--endogenous compound--ec; proteinase --endogenous
compound--ec; matrigel--endogenous compound--ec; cysteine--endogenous compound--ec
 4/3,K/13 (Item 12 from file: 73) Links
    Fulltext available through:
                                       USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
13797099
                EMBASE No: 2006226285
Cysteine cathepsins in the immune response
Zavas(caron)nik-Bergant T.; Turk B.
B. Turk, Department of Biochemistry and Molecular Biology, J. Stefan Institute.
SI-1000 Ljubljana
                        Slovenia
Author Email: boris.turk@ijs.si
Tissue Antigens ( TISSUE ANTIGENS ) ( United Kingdom )
                                                                      2006 , 67/5 (349-355)
                 ISSN: 0001-2815 eISSN: 1399-0039
                                                Page 43
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Document Type: Journal; Review Language: ENGLISH Summa Number Of References: 50 Summary Language: ENGLISH DRUG DESCRIPTORS: * cysteine derivative--endogenous compound--ec; *cathepsin--endogenous compound--ec major histocompatibility antigen class 2--endogenous compound--ec; cathepsin B--endogenous compound--ec; dipeptidyl peptidase I --endogenous compound--ec; cathepsin F--endogenous compound --ec; cathepsin H--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin D--endogenous compound --ec; CD4 antigen--endogenous compound--ec; CD8 antigen--endogenous compound--ec; major histocompatibility antigen class 1--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; cathepsin W --endogenous compound--ec; cathepsin W 4/3,K/14 (Item 13 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2004344716 13702260 Protease expression in interface tissues around loose arthroplasties Kido A.; Pap G.; Nagler D.K.; Ziomek E.; Menard R.; Neumann H.W.; Roessner A. Dr. A. Kido, Department of Orthopedic Surgery, Nara Medical University, 840 Shijo-cho, Kashihara, Nara 634-8522 Japan Author Email: akirakid@naramed-u.ac.jp Clinical Orthopaedics and Related Research (CLIN. ORTHOP. RELAT. RES.) (United 2004 , -/425 (230-236) ISSN: 0009-921X CODEN: CORTB Document Type: Journal; Article Language: ENGLISH Summa Number Of References: 35 Summary Language: ENGLISH DRUG DESCRIPTORS: * proteinase--endogenous compound--ec cathepsin--endogenous compound--ec; interstitial collagenase --endogenous compound --ec; cathepsin B--endogenous compound --ec; cathepsin D--endogenous compound--ec; cathepsin L--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec 4/3,K/15 (Item 14 from file: 73) Links USPTO Full Text Retrieval Options Fulltext available through: **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2006135543 13659084 Lysosomal cysteine proteases: Structure, function and inhibition of cathepsins Prof. R. Roberts, Department of Biology, Program in Biochemistry and Molecular Biology, Ursinus College, P.O. Box 1000, Collegeville, PA 19426-1000 United United States Author Email: rroberts@ursinus.edu Drug News and Perspectives (DRUG NEWS PERSPECT.) (Spain) 2005 , 18/10 (605-614)ISSN: 0214-0934 CODEN: DNPEE Document Type: Journal; Review Language: ENGLISH Summary Language: ENGLISH Number Of References: 111 DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec Tysosome enzyme--endogenous compound--ec; cysteine proteinase --endogenous compound --ec; cathepsin B--endogenous compound --ec; dipeptidyl peptidase Page 44

cathepsinsearch.txt I--endogenous compound--ec; cathepsin F --endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin S--endogenous compound--ec; n [n (3 carboxyoxirane 2 carbonyl)leucyl]agmatine--drug comparison--cm; n [n (3... Drug Terms (Uncontrolled): cathepsin V--endogenous compound--ec; cathepsin O--endogenous compound--ec; cathepsin w--endogenous compound--ec; cathepsin x--endogenous compound--ec; ca 074--drug comparison--cm; ca 074--pharmacology--pd; morpholineurea leucine homophenylalanine vinylsulfonephenyl--drug... 4/3,K/16 (Item 15 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2006036856 13558393 An enzyme-linked immunosorbent assay for human cathepsin X, a potential new inflammatory marker Nagler D.K.; Lechner A.M.; Oettl A.; Kozaczynska K.; Scheuber H.-P.; Gippner-Steppert C.; Bogner V.; Biberthaler P.; Jochum M. D.K. Nagler, Department of Clinical Chemistry and Clinical Biochemistry, University Hospital of Surgery-City, Ludwig-Maximilians-University, Nussbaumstr. 20, 80336 Author Email: dorit.naegler@med.uni-muenchen.de
Journal of Immunological Methods (J. IMMUNOL. METHODS) (Netherlands)
2006 , 308/1-2 (241-250)
CODEN: JIMMB ISSN: 0022-1759 Germany **20 JAN** Publisher Item Identifier: S0022175905003704 Document Type: Journal; Article Summary Language: ENGLISH Language: ENGLISH Number Of References: 35 An enzyme-linked immunosorbent assay for human cathepsin X, a potential new inflammatory marker The human lysosomal cysteine-type carboxypeptidase cathepsin X is mainly present in monocytes and macrophages and may be released into the circulation due. ...inflammatory marker, we have developed a highly sensitive and specific sandwich-type immunoassay (ELISA) for cathepsin X permitting both intra- and extracellular detection and quantification. The dynamic range of the cathepsin X ELISA was determined to be 100 (detection limit) to 8000 pg/ml. Reproducibility of both....of the thiol-dependent cathepsin family was not observed. The ELISA was used to quantify cathepsin X in leukocytes as well as in plasma of healthy volunteers and patients with multiple trauma. During the first 72 h after trauma, plasma levels of cathepsin X increased significantly, particularly in patients who died during the posttraumatic period. In comparison to the well-known inflammation marker neutrophil elastase, cathepsin X levels predicted survival with a higher significance in the later posttraumatic phase. In conclusion, this report provides significance in the later posttraumatic phase. In conclusion, this report provides the first evidence of cathepsin X immunoreactivity not only in cell lysates but also in plasma samples. We suggest that the... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec; elastase--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec 4/3,K/17 (Item 16 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved.

EMBASE No: 2006015689

Endosomal proteases in antigen presentation

13530117

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Chapman H.A.
H.A. Chapman, Department of Medicine, Cardiovascular Research Institute, University of California, San Francisco, CA 94143 United States Author Email: hal.chapman@ucsf.edu
Current Opinion in Immunology ( CURR. OPIN. IMMUNOL. ) ( United Kingdom )
                                                                                                  2006 .
18/1 (78-84)
CODEN: COPIE
                   ISSN: 0952-7915
Publisher Item Identifier: S0952791505002049
Document Type: Journal; Review
                          Summary Language: ENGLISH
Language: ENGLISH
Number Of References: 57
DRUG DESCRIPTORS:
  proteinase--endogenous compound--ec; *major histocompatibility antigen class
1--endogenous compound--ec; *major histocompatibility antigen class 2--endogenous
cathepsin--endogenous compound--ec; cathepsin D--endogenous compound--ec; dipeptidyl
peptidase I--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin S--endogenous compound--ec;
cathepsin L--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
 4/3,K/18 (Item 17 from file: 73) Links
    Fulltext available through:
                                         USPTO Full Text Retrieval Options
EMBASE
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                 EMBASE No: 2005489501
13445732
Gene expression profiles reveal increased mClca3 (Gob5) expression and mucin
production in a murine model of asbestos-induced fibrogenesis
Sabo-Attwood T.; Ramos-Nino M.; Bond J.; Butnor K.J.; Heintz N.; Gruber A.D.; Steele
C.; Taatjes D.J.; Vacek P.; Mossman B.T.
B.T. Mossman, University of Vermont, HSRF 218, 89 Beaumont Ave., Burlington, VT
          United States
Author Email: brooke.mossman@uvm.edu
American Journal of Pathology (AM. J. PATHOL. ) (United States)
                                                                                         2005 . 167/5
(1243 - 1256)
                   ISSN: 0002-9440
CODEN: AJPAA
Document Type: Journal; Article
Language: ENGLISH
                         Summary Language: ENGLISH
Number Of References: 63
DRUG DESCRIPTORS:
* mucin--endogenous compound--ec; *asbestos; *gene product--endogenous compound--ec chrysotile; cyclin B1--endogenous compound--ec; cell cycle protein 20--endogenous
compound--ec; cyclin dependent kinase 1--endogenous compound--ec; chemokine--endogenous compound--ec; complement component C1--endogenous compound--ec; chitinase--endogenous compound--ec; tumor necrosis factor derivative--endogenous compound--ec; interleukin 1beta--endogenous compound--ec;
macrophage elastase--endogenous compound--ec; stromelysin --endogenous compound--ec;
integrin--endogenous compound--ec; cathepsin K--endogenous compound--ec;
cathepsin--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin
S--endogenous compound--ec; cytokine--endogenous compound--ec; unindexed drug;
unclassified drug
Drug Terms (Uncontrolled): protein mCLCA3--endogenous compound--ec; protein
Gob5--endogenous compound--ec; CDC28 protein kinase regulatory subunit 2--endogenous compound--ec; CCL9 chemokine--endogenous compound--ec; ccl6 chemokine--endogenous compound--ec; chitinase 3 like 3--endogenous compound--ec; tumor necrosis factor
superfamily member 10 --endogenous compound--ec; integrin alphax--endogenous
compound--ec; cathepsin Z--endogenous compound --ec
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4/3,K/19 (Item 18 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options Page 46

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13365751
                     EMBASE No: 2005432915
Large scale real-time PCR analysis of mRNA abundance in rainbow trout eggs in
relationship with egg quality and post-ovulatory ageing
Aegerter S.; Jalabert B.; Bobe J.
J. Bobe, INRA, SCRIBE, Campus de Beaulieu, F-35042 Rennes Cedex Author Email: Julien.Bobe@rennes.inra.fr
Molecular Reproduction and Development (MOL. REPROD. DEV.) (United States) 2005, 72/3 (377-385) CODEN: MREDE ISSN: 1040-452X
Document Type: Journal; Article
Language: ÉNGLISH
                              Summary Language: ENGLISH
Number Of References: 35
...period, eight transcripts (nucleoplasmin or Npm2, ferritin H, tubulin beta, JNK1,
cyclin A1, cyclin A2, cathepsin Z, and IGF2) exhibited a differential abundance at one or several collection time(s). Interestingly, we....lower levels of Npm2, tubulin beta, and IGF1 transcripts. In contrast, keratins 8 and 18, cathepsin Z, and prostaglandin synthase 2 were more abundant in low quality eggs than in high
quality..
DRUG DESCRIPTORS:
nucleoplasmin--endogenous compound--ec; somatomedin--endogenous compound--ec; cyclin
A--endogenous compound--ec; tubulin --endogenous compound--ec; ferritin--endogenous
compound--ec; stress activated protein kinase 1--endogenous compound--ec; cathepsin--endogenous compound--ec; somatomedin C--endogenous compound--ec; beta tubulin--endogenous compound--ec; keratin --endogenous compound--ec; prostaglandin synthase--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cyclin al--endogenous compound--ec; cyclin A2--endogenous compound--ec; cathepsin Z--endogenous compound --ec; keratin 8--endogenous compound--ec; keratin 18--endogenous compound--ec; prostaglandin synthase 2--endogenous compound --ec
  4/3,K/20 (Item 19 from file: 73) Links
     Fulltext available through:
                                                     USPTO Full Text Retrieval Options
(c) 2007 Elsevier B.V. All rights reserved.
                     EMBASE No: 2005386832
13318701
Up-regulation of cathepsin X in Helicobacter pylori gastritis and gastric cancer
Krueger S.; Kalinski T.; Hundertmark T.; Wex T.; Kuster D.; Peitz U.; Ebert M.; Nagler D.K.; Kellner U.; Malfertheiner P.; Naumann M.; Rocken C.; Roessner A. S. Krueger, Department of Pathology, Otto-von-Guericke University, Leipziger Strasse
44. D-39120 Magdeburg
                                    Germany
Author Email: Sabine.Krueger@medizin.uni-magdeburg.de
Journal of Pathology ( J. PATHOL. ) ( United Kingdom ) CODEN: JPTLA ISSN: 0022-3417
                                                                                           2005 , 207/1 (32-42)
CODEN: JPTLA
Document Type: Journal; Article
Language: ENGLISH Summ
Number Of References: 35
                                Summary Language: ENGLISH
Up-regulation of cathepsin X in Helicobacter pylori gastritis and gastric cancer
Recently, we identified increased cathepsin X expression in H. pylori-infected
gastric mucosa. Here, we describe further up-regulation in gastric cancer and report
on the role of inflammatory cytokines required for cathepsin X up-regulation in H.
pylori-infected gastric mucosa, as well as on consequences for cellular...
...infected and non-infected patients. Gastric cancer samples were obtained from
patients undergoing gastric surgery. Cathepsin X was detected in gastric mucosa by quantitative real-time RT-PCR, western blotting and immunohistochemistry. Induction of cathepsin X expression in epithelial and inflammatory cells caused by H. pylori infection was tested in in.....cultures of AGS cells and monocytic cells. Patients
                                                              Page 47
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with H. pylori gastritis showed significantly higher cathepsin X mRNA (2.5-fold) and protein (1.6-fold) expression than H. pylori-negative patients. Cathepsin X was also up-regulated in gastric cancer (3-12-fold) compared to non-neoplastic mucosa. Cathepsin X was predominantly expressed by macrophages in the mucosal stroma and in glands of the antral mucosa. In addition, tumour cells stained for cathepsin X in 26 (68%) patients with gastric carcinoma. In general, staining was significantly more common (20....via soluble factors in the culture medium seems to be responsible for increased expression of cathepsin X in monocytes. Using antisense oligonucleotides, cathepsin X up-regulation was directly associated with higher invasiveness in vitro. Although no correlation of cathepsin X expression and TNM stage was found, our study demonstrates that cathepsin X plays a role not only in the chronic inflammation of gastric mucosa but also in...

DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec

cytokine--endogenous compound--ec; messenger RNA--endogenous compound--ec; antisense oligonucleotide; unclassified drug

Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/21 (Item 20 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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13163079 EMBASE No: 2005224520
Carboxypeptidases cathepsins X and B display distinct protein profile in human cells

and tissues

Kos J.; Sekirnik A.; Premzl A.; Bergant V.Z.; Langerholc T.; Turk B.; Werle B.; Golouh R.; Repnik U.; Jeras M.; Turk V.
J. Kos, Department of Pharmaceutical Biology, Faculty of Pharmacy, University of Ljubljana, As(caron)kerc(caron)eva 7, SI-1000 Ljubljana Slovenia
Author Email: janko.kos@krka.biz
Experimental Cell Research (EXP. CELL RES.) (United States) 15 MAY 2005, 306/1 (103-113)
CODEN: ECREA ISSN: 0014-4827
Publisher Item Identifier: S0014482704007220
Document Type: Journal; Article
Language: ENGLISH Summary Language: ENGLISH
Number Of References: 42

Cathepsin X, a recently discovered lysosomal cysteine protease, shares common structural features and activity properties with cysteine.....distribution in cells and tissues and to their possible roles in malignancy. Protein level of cathepsin X did not differ significantly between matched pairs of lung tumor and adjacent lung tissue obtained......6-fold higher in tumor compared to adjacent lung tissue. Immunohistochemical analysis of lung tumor cathepsin X revealed very faint staining in tumor cells but positive staining in infiltrated histiocytes, alveolar macrophages, bronchial epithelial cells, and alveolar type II cells. Cathepsin X stained positive also in CD68SUP+ cells in germinal centers of secondary follicles in lymph nodes10A neoT and MDA-MB 231, showed positive staining for cathepsin B, but negative for cathepsin X. We showed that the invasive potential of MCF-10A neoT cells can be impaired by specific inhibitor of cathepsin B but not by that of cathepsin X. Cathepsin X was found in large amounts in the pro-monocytic U-937 cell line, in monocytes and in dendritic cells, generated from monocytes in vitro. Our results show that cathepsin X is not involved in degradation of extracellular matrix, a proteolytic event leading to tumor cell...

DRUG DESCRIPTORS:

* carboxypeptidase--endogenous compound--ec; *cathepsin B--endogenous compound--ec CD68 antigen--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/22 (Item 21 from file: 73) Links

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Fulltext available through:
                                         USPTO Full Text Retrieval Options
EMBASE
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13140612
                EMBASE No: 2005207535
Capturing protein interactions in the secretory pathway of living cells
Nyfeler B.; Michnick S.W.; Hauri H.-P.
H.-P. Hauri, Dept. of Pharmacol. and Neurbiology, Biozentrum, University of Basel,
Klingelbergstrasse 70, CH-4056 Basel
                                                 Switzerland
Author Email: hans-peter.hauri@unibas.ch
Proceedings of the National Academy of Sciences of the United States of America (PROC. NATL. ACAD. SCI. U. S. A.) (United States) 03 MAY 2005, 102/18
(6350 - 6355)
                   ISSN: 0027-8424
CODEN: PNASA
Document Type: Journal; Article
Language: ENGLISH
                        Summary Language: ENGLISH
Number Of References: 35
...ERGIC-53-interacting multicoagulation factor deficiency protein MCFD2, and to
ERGIC-53's cargo glycoprotein cathepsin Z. YFP PCA analysis revealed the
oligomerization of ERGIC-53 and its interaction with MCFD2, as well as its lectin-mediated interaction with cathepsin Z. Mutation of the lectin domain of ERGIC-53 selectively decreased YFP complementation with cathepsin Z. Using YFP PCA,
we discovered a carbohydrate-mediated interaction between ERGIC-53 and cathepsin
С..
DRUG DESCRIPTORS:
  endoplasmic reticulum golgi intermediate compartment protein 53--endogenous
compound--ec; *cathepsin--endogenous compound--ec; *hybrid protein--endogenous
compound--ec
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; multicoaqulation
factor deficiency protein--endogenous compound--ec
 4/3,K/23 (Item 22 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
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                EMBASE No: 2005198390
The human brain mannose 6-phosphate glycoproteome: A complex mixture composed of
multiple isoforms of many soluble lysosomal proteins
Sleat D.E.; Lackland H.; Wang Y.; Sohar I.; Xiao G.; Li H.; Lobel P. Dr. P. Lobel, Ctr. for Adv. Biotech. and Medicine, 679 Hoes Lane, Piscataway, NJ
          United States
Author Email: lobel@cabm.rutgers.edu
Proteomics ( PROTEOMICS ) ( Germany )
                                                   2005 , 5/6 (1520-1532)
CODEN: PROTC
                 ISSN: 1615-9853
Document Type: Journal ; Article
                         Summary Language: ENGLISH
Language: ENGLISH
Number Of References: 41
DRUG DESCRIPTORS:
* mannose 6 phosphate--endogenous compound--ec; *lysosome enzyme --endogenous
compound--ec; *proteome--endogenous compound--ec somatomedin B receptor--endogenous compound--ec; n acetyl beta
glucosaminidase--endogenous compound--ec; cathepsin S--endogenous compound--ec; deoxyribonuclease II--endogenous compound--ec; dipeptidyl peptidase--endogenous
compound--ec; gamma glutamyl hydrolase--endogenous compound--ec;
legumain--endogenous compound--ec; lysophospholipase--endogenous compound--ec;
proline carboxypeptidase--endogenous compound--ec; clusterin --endogenous
compound--ec; acetylesterase--endogenous compound--ec; alpha mannosidase--endogenous compound--ec; serine carboxypeptidase--endogenous compound--ec; ribonuclease --endogenous compound--ec; ependymin--endogenous compound--ec; n4 (beta n acetylglucosaminyl)asparaginase--endogenous compound --ec;
                                                 Page 49
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angiotensinogen--endogenous compound--ec; cerebroside sulfatase--endogenous compound--ec; acylsphingosine deacylase --endogenous compound--ec; palmitoyl protein thioesterase --endogenous compound--ec; cystatin C--endogenous compound --ec; cystatin B--endogenous compound--ec; dipeptidyl peptidase I --endogenous
compound--ec; cathepsin D--endogenous compound --ec; cathepsin F--endogenous
compound--ec; cathepsin L--endogenous compound--ec; cathepsin--endogenous
compound--ec; F box protein--endogenous compound--ec; ferritin--endogenous
compound--ec; alpha levo fucosidase--endogenous compound--ec; alpha
glucosidase--endogenous compound--ec; sulfatase--endogenous compound--ec; alpha
galactosidase--endogenous compound--ec; beta galactosidase--endogenous compound--ec; beta glucuronidase --endogenous compound--ec; beta n acetylhexosaminidase
A--endogenous compound--ec; beta n acetylhexosaminidase B--endogenous compound--ec; iduronate 2 sulfatase--endogenous compound--ec; levo iduronidase--endogenous compound--ec; galectin 1--endogenous compound--ec; acid lipase--endogenous
compound--ec; myelin associated glycoprotein--endogenous compound--ec; beta
mannosidase --endogenous compound--ec; prosaposin--endogenous compound --ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; cathepsin
p--endogenous compound--ec; cathepsin x --endogenous compound--ec; dipeptidyl
peptidase VII--endogenous compound--ec; ribonuclease 6--endogenous compound--ec; n
acetyl 6 galactosamine sulfatase--endogenous compound--ec; n acetyl glucosamine 6
sulfatase--endogenous compound--ec
  4/3,K/24 (Item 23 from file: 73) Links
                                                USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
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                   EMBASE No: 2005198358
Gene expression profiling of the effect of high-dose intravenous Ig in patients with
Kawasaki disease
Abe J.; Jibiki T.; Noma S.; Nakajima T.; Saito H.; Terai M. Dr. J. Abe, Department of Allergy and Immunology, Natl. Res. Inst. Child Hlth./Devmt., 2-10-1 Ohkura, Setagaya-ku, Tokyo 157-8535 Japan Author Email: jabe@nch.go.jp
Journal of Immunology (J. IMMUNOL.) (United States)
                                                                                     01 MAY 2005 . 174/9
(5837 - 5845)
CODEN: JOIMA
                      ISSN: 0022-1767
Document Type: Journal; Article
Language: ENGLISH
                             Summary Language: ENGLISH
Number Of References: 53
DRUG DESCRIPTORS:

* ...dose--do; *immunoglobulin--drug therapy--dt; *immunoglobulin --intravenous drug administration--iv; *chemokine receptor CCR2--endogenous compound--ec; *protein S
100--endogenous compound--ec; *Fc receptor--endogenous compound--ec;
*adrenomedullin--endogenous compound--ec
formylpeptide receptor--endogenous compound--ec; C reactive protein --endogenous
compound--ec; toll like receptor 2--endogenous compound--ec; adiponectin--endogenous
compound--ec; cell surface receptor--endogenous compound--ec; colony stimulating
factor receptor--endogenous compound--ec; interleukin 8 receptor --endogenous
compound--ec; CD39 antigen--endogenous compound --ec; CD16 antigen--endogenous compound--ec; colony stimulating factor 1--endogenous compound--ec; protein tyrosine
phosphatase --endogenous compound--ec; protein p57--endogenous compound--ec; interleukin 3--endogenous compound--ec; versican--endogenous compound--ec; immunoglobulin kappa chain--endogenous compound --ec; APRIL protein--endogenous compound--ec; dysferlin--endogenous compound--ec; chimerin--endogenous compound--ec; hematopoietic cell kinase--endogenous compound--ec; phosphatase --endogenous
compound--ec; RGS2 protein--endogenous compound --ec; Rab protein--endogenous
compound--ec; transcription factor --endogenous compound--ec; protein v fos--endogenous compound --ec; early growth response factor 1--endogenous
compound--ec; calreticulin--endogenous compound--ec; major histocompatibility antigen class 2--endogenous compound--ec; hexokinase--endogenous compound--ec; 5
                                                         Page 50
```

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aminolevulinate synthase--endogenous compound --ec; oxidoreductase--endogenous compound--ec; cytochrome P450 1B1 --endogenous compound--ec; long chain fatty acid
coenzyme A ligase --endogenous compound--ec; histidine ammonialyase--endogenous
compound--ec; microsomal aminopeptidase--endogenous compound --ec; spermidine--endogenous compound--ec; acyltransferase --endogenous compound--ec;
cathepsin--endogenous compound--ec; collapsin response mediator protein--endogenous
compound--ec; ribosome protein--endogenous compound--ec; aquaporin 9--endogenous
compound--ec; carrier protein--endogenous compound--ec; scramblase--endogenous
compound-ec; heat shock protein 70 --endogenous compound-ec; protein disulfide isomerase--endogenous compound-ec; unclassified drug
Drug Terms (Uncontrolled): protein S100A9--endogenous compound-ec; protein
$100A12--endogenous compound--ec; protein $100A8--endogenous compound--ec; adiponectin receptor 1--endogenous compound--ec; leukocyte immunoglobulin receptor B1--endogenous compound--ec; leukocyte immunoglobulin receptor B2--endogenous compound--ec; leukocyte immunoglobulin like receptor B3--endogenous compound--ec;
stabilin 1 --endogenous compound--ec; S phase response protein--endogenous
compound--ec; growth arrest specific protein 7--endogenous compound--ec; cold autoinflammatory syndrome 1 protein--endogenous compound--ec; pre B cell colony
enhancing factor--endogenous compound--ec; proapoptotic caspase adaptor protein--endogenous compound--ec; chimerin 2--endogenous compound--ec; dual
specificity phosphatase 1--endogenous compound--ec; Rab31 protein --endogenous compound--ec; kruppel like factor 4--endogenous compound--ec; cold shock domain protein A--endogenous compound--ec; SFFV proviral integration 1 protein--endogenous compound--ec; transcription factor 7 like 2--endogenous compound--ec; hexokinase 3--endogenous compound--ec; guanosine phosphate reductase--endogenous compound--ec;
biliverdin reductase B--endogenous compound--ec; gamma interferon inducible protein
30--endogenous compound--ec; flavoprotein oxidoreductase --endogenous compound--ec; neutrophil cytosolic factor 2--endogenous compound--ec; spermine n1
acetyltransferase--endogenous compound--ec; cathepsin Z--endogenous compound --ec;
mitochondrial solute carrier protein--endogenous compound--ec
```

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4/3,K/25 (Item 24 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
(c) 2007 Elsevier B.V. All rights reserved.
                 EMBASE No: 2005147588
13086504
Pancreatic beta-cell failure and diabetes in mice with a deletion mutation of the
endoplasmic reticulum molecular chaperone gene P58SUPIPK
Ladiges W.C.; Knoblaugh S.E.; Morton J.F.; Korth M.J.; Sopher B.L.; Baskin C.R.;
MacAuley A.; Goodman A.G.; LeBoeuf R.C.; Katze M.G.
W.C. Ladiges, Department of Comparative Medicine, Box 357190, University of
Washington, Seattle, WA 98195 United S
Author Email: wladiges@u.washington.edu
                                          United States
Diabetes ( DIABETES ) ( United States )
                                                        2005 , 54/4 (1074-1081)
                  ISSN: 0012-1797
CODEN: DIAEA
Document Type: Journal; Article
                          Summary Language: ENGLISH
Language: ENGLISH
Number Of References: 26
DRUG DESCRIPTORS:
* chaperone--endogenous compound--ec; *protein p58--endogenous compound--ec
glucose; initiation factor 2alpha-endogenous compound-ec; cathepsin L-endogenous compound-ec; protein p53-endogenous compound-ec; lymphotoxin beta-endogenous compound-ec; cathepsin D-endogenous compound-ec; cathepsin B-endogenous compound-ec; serine proteinase Omi-endogenous compound-ec; FAS ligand-endogenous
compound--ec; cathepsin--endogenous compound--ec; annexin--endogenous compound--ec;
cytochrome c --endogenous compound--ec; STAT3 protein--endogenous compound --ec; beta arrestin--endogenous compound--ec; immunoglobulin enhancer binding
protein--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec
```

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cathepsinsearch.txt
 4/3,K/26 (Item 25 from file: 73) Links
                                      USPTO Full Text Retrieval Options
   Fulltext available through:
(c) 2007 Elsevier B.V. All rights reserved.
               EMBASE No: 2004261313
12666149
Cathepsins K, L, B, X and W are differentially expressed in normal and chronically
inflamed gastric mucosa
Buhling F.; Peitz U.; Kruger S.; Kuster D.; Vieth M.; Gebert I.; Roessner A.; Weber
E.; Malfertheiner P.; Wex T.
T. Wex, Dept. of Gastroenterology, Dept. of Infectious Disease, Leipziger Str. 44.
D-39120 Magdeburg
                      Germany
Author Email: thomas.wex@medizin.uni-magdeburg.de
Biological Chemistry (BIOL. CHEM.) (Ğermany)
CODEN: BICHF ISSN: 1431-6730
                                                          2004 , 385/5 (439-445)
Document Type: Journal; Article
Language: ENGLISH Summanumber Of References: 28
                     Summary Language: ENGLISH
...was expressed at very low levels. Infection by Helicobacter pylori caused a significant induction of cathepsin X (p<0.008), whereas the other cathepsins were
not or only locally affected by H. pylori infection or reflux disease.
Immunohistochemistry revealed specific expression of cathepsin X (macrophages),
cathepsin K (parietal cells) and cathepsin W (lymphocytes), whereas cathepsins B and
L were..
DRUG DESCRIPTORS:
* cathepsin K--endogenous compound--ec; *cathepsin L--endogenous compound--ec; *cathepsin B--endogenous compound--ec; * cathepsin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; cathepsin w
--endogenous compound--ec
 4/3,K/27 (Item 26 from file: 73) Links
   Fulltext available through:
                                     USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
               EMBASE No: 2004255824
Up-regulation of cathepsin X in prostate cancer and prostatic intraepithelial
neoplasia
Nagler D.K.; Kruger S.; Kellner A.; Ziomek E.; Menard R.; Buhtz P.; Krams M.; Roessner A.; Kellner U.
D.K. Nagler, Dept. of Clin. Chem./Clin. Biochem., University Hospital of Surgery-City, Ludwig-Maximilians-University, Nussbaumstr. 20, 80336 Munich
                                                                                        Germany
Author Email: dorit.naegler@clinbio.med.uni-muenchen.de
                                                01 JUL 2004 , 60/2 (109-119)
Prostate ( PROSTATE ) ( United States )
                ISSN: 0270-4137
CODEN: PRSTD
Document Type: Journal; Article
Language: ENGLISH Summa
Number Of References: 36
                      Summary Language: ENGLISH
Up-regulation of cathepsin X in prostate cancer and prostatic intraepithelial
neoplasia
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec
cathepsin F--endogenous compound--ec; cathepsin B--endogenous compound--ec;
cathepsin L--endogenous compound--ec; genomic DNA--endogenous compound--ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
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4/3,K/28 (Item 27 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options
Page 52

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(c) 2007 Elsevier B.V. All rights reserved.
                    EMBASE No: 2004197020
Human B lymphoblastoid cells contain distinct patterns of cathepsin activity in
endocytic compartments and regulate MHC class II transport in a cathepsin
S-independent manner
Lautwein A.; Kraus M.; Reich M.; Burster T.; Brandenburg J.; Overkleeft H.S.;
Schwarz G.; Kammer W.; Weber E.; Kalbacher H.; Nordheim A.; Driessen C.
C. Driessen, MNF Universitat Tubingen, Ob dein Himmelreich 7, 72074 Tubingen
Author Email: christoph.driessen@med.uni-tuebingen.de
Journal of Leukocyte Biology ( J. LEUKOCYTE BIOL. ) ( United States )
                                                                                                                2004 , 75/5
 (844 - 855)
CODEN: JLBIE
                       ISSN: 0741-5400
Document Type: Journal; Article
Language: ENGLISH
                               Summary Language: ENGLISH
Number Of References: 56
DRUG DESCRIPTORS:
* major histocompatibility antigen class 2--endogenous compound--ec; * cathepsin--endogenous compound--ec; *cathepsin S--endogenous compound--ec proteinase--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin D--endogenous compound--ec; dipeptidyl peptidase I--endogenous compound--ec; HLA DM antigen; vinyl derivative; phenol
derivative; unclassified drug
Drug Terms (Uncontrolled): asparagine specific endoprotease--endogenous compound--ec; cathepsin Z; leucine homophenylalanine vinylsulfone phenol
  4/3,K/29 (Item 28 from file: 73) Links
     Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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 12535984
                    EMBASE No: 2004129918
Expression and characterization of cathepsin P
Mason R.W.; Bergman C.A.; Lu G.; Frenck Holbrook J.; Sol-Church K.
R.W. Mason, Department of Biomedical Research, Alfred I. duPont Hosp. for Children,
1600 Rockland Road, Wilmington, DE 19803
                                                                   United States
Author Email: mason@medsci.udel.edu
Biochemical Journal (BIOCHEM. J.) (United Kingdom) 01 MAR 2004, 378/2
(657-663)
                       ISSN: 0264-6021
CODEN: BIJOA
Document Type: Journal; Article
Language: ENGLISH Summ
Number Of References: 24
                             Summary Language: ENGLISH
Expression and characterization of cathepsin P
 ...in placental tissues of all mammalian species. In the present study, it was shown
that cathepsin P can be expressed in Pichia pastoris as an inactive zymogen that can
be activated with proteinase K, chymotrypsin or pancreatic elastase at neutral pH.
Unlike other mammalian cathepsins, cathepsin P could also be autoactivated at
neutral pH, but not at acidic pH. The activated enzyme....SUB2SOSUB4 and
hyaluronate stimulated the activity of the protease against peptidyl substrates. The
properties of cathepsin P appear to be quite distinct from those of cathepsin L,
indicating that the duplication that gave rise to cathepsin P has probably not
yielded an enzyme that provides a subfunction of cathepsin L in rodents. It seems
probable that cathepsin P has evolved to perform a function that is performed by an
probable that cathepsin P has evolved to perform a function that is performed by an
evolutionarily unrelated protease in...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cathepsin L--endogenous compound--ec
messenger RNA--endogenous compound--ec; proteinase K; chymotrypsin; pancreatic elastase; peptide derivative--endogenous compound--ec; protein
                                                            Page 53
```

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cathepsinsearch.txt
derivative--endogenous compound--ec; transferrin--endogenous compound--ec; inorganic
salt; sodium sulfate; hyaluronic acid; unclassified drug
Drug Terms (Uncontrolled): cathepsin p--endogenous compound--ec; azocasein
--endogenous compound--ec
  4/3,K/30 (Item 29 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
12534119
                 EMBASE No: 2004127448
Cysteine proteases as disease markers
Berdowska I.
I. Berdowska, Department of Medical Biochemistry, Wroclaw Medical University, 10
Chalubinskiego, 50-368 Wroclaw
                                          Poland
Author Email: iza@bioch.am.wroc.pl
Clinica Chimica Acta ( CLIN. CHIM. ACTA ) ( Netherlands ) 2004 , 342/1-2 (41-69) CODEN: CCATA ISSN: 0009-8981
Publisher Item Identifier: S0009898103006041
Document Type: Journal; Review
Language: ENGLISH Summar
Number Of References: 248
                        Summary Language: ENGLISH
DRUG DESCRIPTORS:
* cysteine proteinase--endogenous compound--ec; *tumor marker --endogenous
compound--ec
peptidase--endogenous compound--ec; papain--endogenous compound--ec; cysteine
derivative--endogenous compound--ec; cathepsin B--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin S--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin F--endogenous compound--ec; dipeptidyl peptidase I --endogenous compound--ec; protein precursor--endogenous compound--ec; enzyme precursor--endogenous compound--ec; hormone precursor--endogenous compound--ec; major histocompatibility antigen class 2--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cysteine cathepsin derivative--endogenous compound--ec;
cathepsin v --endogenous compound--ec; cathepsin x--endogenous compound--ec;
cathepsin w--endogenous compound--ec; cathepsin o--endogenous compound--ec
 4/3,K/31 (Item 30 from file: 73) Links
                                         USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
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                EMBASE No: 2004098453
12503843
Myxobolus cerebralis: Identification of a cathepsin Z-like protease gene (MyxCP-1)
expressed during parasite development in rainbow trout, Oncorhynchus mykiss
Kelley G.O.; Adkison M.A.; Leutenegger C.M.; Hedrick R.P.
G.O. Kelley, Dept. of Medicine and Epidemiology, School of Veterinary Medicine,
University of California, Davis, CA 95616
                                                       United States
Author Email: gokelley@ucdavis.edu
Experimental Parasitology (EXP. PARASITOL.) (United States) 2003, 105/3-4
 (201-210)
                   ISSN: 0014-4894
CODEN: EXPAA
Document Type: Journal; Article
Language: ENGLISH
                         Summary Language: ENGLISH
Number Of References: 54
Myxobolus cerebralis: Identification of a cathepsin Z-like protease gene (MyxCP-1)
expressed during parasite development in rainbow trout, Oncorhynchus mykiss
...cysteine proteases. MyxCP-1 features a propeptide region and sequence insertions that are characteristics of cathepsin Z proteases. Phylogenetic comparisons of M.
cerebralis to other eukaryotes based on full-length cathepsin-like genes show that
                                                 Page 54
```

MyxCP-1 is the earliest lineage in the cathepsin Z group and separated from cathepsin L, B, and C-like proteases. Using TagMan PCR differential levels of transcription of the cathepsin Z-like protease were found in earlier and later developmental stages of the parasite in experimentally... DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin Z like protease--endogenous compound--ec

4/3,K/32 (Item 31 from file: 73) Links

USPTO Full Text Retrieval Options Fulltext available through:

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EMBASE No: 2004079961 12487347

The Caenorhabditis elegans Cathepsin Z-like Cysteine Protease, Ce-CPZ-1, Has a Multifunctional Role during the Worms' Development

Hashmi S.; Zhang J.; Oksov Y.; Lustigman S.

S. Hashmi, Laboratory of Molecular Parasitology, Lindsley F. Kimball Research Inst., New York Blood Center, 310 E. 67th St., New York, NY 10021 United States

Author Email: shashmi@nybloodcenter.org

Journal of Biological Chemistry (J. BIOL. CHEM.) (United States) 279/7 (6035-6045)
CODEN: JBCHA ISSN: 0021-9258 13 FEB 2004 .

Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 60

The Caenorhabditis elegans Cathepsin Z-like Cysteine Protease, Ce-CPZ-1, Has a Multifunctional Role during the Worms' Development

We have analyzed the expression and function of Ce-cpz-1, a Caenorhabditis elegans cathepsin Z-like cysteine protease gene, during development of the worm. The cpz-1 gene is expressed....are degraded prior to shedding and ecdysis. The similar localization of the related Onchocerca volvulus cathepsin Z protein suggests that the function of CPZ-1 during molting might be conserved in other....basement membrane extracellular matrix assembly process. The present findings have defined a critical role for cathepsin Z in nematode biology. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; cathepsin Z like cysteine proteinase--endogenous compound--ec

4/3,K/33 (Item 32 from file: 73) Links

Fulltext available through: USPTO Full Text Retrieval Options **EMBASE**

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EMBASE No: 2004009245 12409511

Identification of differentially expressed genes in models of melanoma progression by cDNA array analysis: SPARC, MIF and a novel cathepsin protease characterize aggressive phenotypes

Rumpler G.; Becker B.; Hafner C.; McClelland M.; Stolz W.; Landthaler M.; Schmitt R.; Bosserhoff A.; Vogt T.

Dr. T. Vogt, Department of Dermatology, University of Regensburg, D-93042 Regensburg

Author Email: thomas.voqt@klinik.uni-regensburg.de

Experimental Dermatology (EXP. DERMATOL.) (United Kingdom) $2003 \cdot 12/6$

(761-771)

CODEN: EXDEE ISSN: 0906-6705 Document Type: Journal; Article Language: ENGLISH Summary Language

Summary Language: ENGLISH

Number Of References: 46

...migration inhibiting factor (MIF), an important modulator of both cell cycle progression and angiogenesis, and cathepsin Z, a novel member of the family of matrix degrading proteinases. (c) Blackwell Munksgaard, 2003. DRUG DESCRIPTORS: * complementary DNA--endogenous compound--ec; *osteonectin--endogenous compound--ec: *macrophage migration inhibition factor--endogenous compound--ec; *cathepsin--endogenous compound--ec; * proteinase--endogenous compound--ec reduced nicotinamide adenine dinucleotide dehydrogenase (ubiquinone) --endogenous compound--ec; ubiquitin--endogenous compound--ec; selenoprotein--endogenous compound--ec; tumor protein--endogenous compound--ec; guanine nucleotide binding protein--endogenous compound--ec; HLA antigen class 2--endogenous compound--ec; laminin binding protein--endogenous compound--ec; protein --endogenous compound--ec; polyadenylic acid binding protein --endogenous compound--ec; DNA binding protein--endogenous compound--ec; polyadenylic acid binding protein --endogenous compound--ec; polyadenylic acid binding protein --endogenous compound--ec; polyadenylic acid binding protein --endogenous compound--ec; polyadenylic acid binding protein--endogenous compound--ec; polyadenylic acid binding protein --endogenous compound--ec; polyadenylic acid binding protei shock protein 90--endogenous compound--ec; cytochrome b --endogenous compound--ec; protein lysine 6 oxidase--endogenous compound--ec; adenosine triphosphatase--endogenous compound --ec; receptor--endogenous compound--ec; phospholipid transfer protein--endogenous compound--ec; beta galactosidase--endogenous compound--ec; unindexed drug; unclassified drug Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; Wilm tumor related protein--endogenous compound--ec; guanine nucleotide binding protein beta subunit like protein--endogenous compound--ec; eukaryotic translation elongation factor 1 gamma--endogenous compound--ec; glia derived nexin--endogenous compound--ec; folic acid receptor 1--endogenous compound--ec 4/3,K/34 (Item 33 from file: 73) Links USPTO Full Text Retrieval Options Fulltext available through: (c) 2007 Elsevier B.V. All rights reserved. 12290413 EMBASE No: 2003402858 Phylogeny of antigen-processing enzymes: Cathepsins of a cephalochordate, an agnathan and a bony fish Uinuk-Ool T.S.; Takezaki N.; Kuroda N.; Figueroa F.; Sato A.; Samonte I.E.; Mayer W.E.; Klein J. T.S. Uinuk-Ool, Max-Planck-Inst. fur Biologie, Abteilung Immungenetik, Corrensstrasse 42, D-72076 Tubingen Author Email: tanya@tuebingen.mpg.de Scandinavian Journal of Immunology (SCAND. J. IMMUNOL.) (United Kingdom) 01 OCT 2003 , 58/4 (436-448) CODEN: SJIMA ISSN: 0300-9475 Document Type: Journal; Article Language: ENGLISH Summa Number Of References: 71 Summary Language: ENGLISH DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec cathepsin B--endogenous compound--ec; cathepsin L--endogenous compound--ec; cathepsin F--endogenous compound--ec; dipeptidyl peptidase I--endogenous compound--ec; cathepsin S --endogenous compound--ec; cathepsin K--endogenous compound --ec; complementary DNA; unclassified drug Drug Terms (Uncontrolled): cathepsin Z--endogenous compound--ec; cathepsin O --endogenous compound--ec 4/3,K/35 (Item 34 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2003014131 11901450 Determination of the mRNA sequence of cathepsin Y, a cysteine endopeptidase from rat spleen, and confirmation of its ubiquitous expression ${\sf P}$ Page 56

```
Nakazono E.; Kamata Y.; Yamafuji K.
K. Yamafuji, Division of Food and Nutrition, Nakamura Gakuen University, Befu 5-7-1,
Jonan-ku, Fukuoka 814-0198
                                    Japan
Biological Chemistry (BIOL. CHEM.) (Germany)
CODEN: BICHF ISSN: 1431-6730
                                                                01 DEC 2002 , 383/12 (1971-1975)
Document Type: Journal; Article
Language: ENGLISH
                         Summary Language: ENGLISH
Number of References: 14
Determination of the mRNA sequence of cathepsin Y, a cysteine endopeptidase from rat
spleen, and confirmation of its ubiquitous expression
...by its action of producing kinin-potentiating peptide from a plasma protein. We named it cathepsin Y due to its localization, acidic pH optimum and the presence of the same set of.....the mRNA sequence resulted in the omission of the strangely attached C-terminal peptide from cathepsin Y.
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
messenger RNA--endogenous compound--ec; cysteine proteinase --endogenous
compound--ec; amino acid--endogenous compound --ec; thiol--endogenous compound--ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin Y--endogenous compound--ec
 4/3,K/36 (Item 35 from file: 73) Links
                                         USPTO Full Text Retrieval Options
    Fulltext available through:
EMBASE
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                EMBASE No: 2001324286
11309979
Lysosomal cysteine proteases: Facts and opportunities
Turk V.; Turk B.; Turk D.
V. Turk, Department of Biochemistry, J. Stefan Institute, Ljubljana
Author Email: vito.turk@ijs.si
EMBO Journal (EMBO J.) (United Kingdom) 03 SEP 2001, 20/17 (4629-4633)
CODEN: EMJOD ISSN: 0261-4189
Document Type: Journal; Review
Language: ENGLISH
                         Summary Language: ENGLISH
Number Of References: 45
DRUG DESCRIPTORS:
* cysteine proteinase--endogenous compound--ec; *cathepsin--endogenous compound--ec
enzyme precursor--endogenous compound--ec; amino acid; cathepsin L --endogenous
compound--ec; cathepsin S--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin H--endogenous compound--ec; unclassified drug
Drug Terms (Uncontrolled): cathepsin V--endogenous compound--ec; cathepsin
w--endogenous compound--ec; cathepsin o--endogenous compound--ec; cathepsin
x--endogenous compound--ec
 4/3,K/37 (Item 36 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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11229615
                EMBASE No: 2001244434
Cathepsins X and B display distinct activity profiles that can be exploited for
inhibitor design
Menard R.; Therrien C.; Lachance P.; Sulea T.; Qi H.; Alvarez-Hernandez A.; Roush
R. Menard, Biotechnology Research Institute, National Research Council of Canada, 6100 Royalmount Avenue, Montreal, Que. H4P 2R2 Canada
                                                 Page 57
```

```
cathepsinsearch.txt
Biological Chemistry (BIOL. CHEM.) (Germany)
CODEN: BICHF ISSN: 1431-6730
                                                                        2001 , 382/5 (839-845)
Document Type: Journal ; Article
Language: ENGLISH Summary Language: ENGLISH
Number Of References: 21
  ..share similar activity profiles against substrates and inhibitors. Using quenched
fluorogenic substrates, we show that cathepsin X preferentially cleaves substrates
through a monopeptidyl carboxypeptidase pathway, while cathepsin B displays a
preference for....approximately 2 orders of magnitude. Cleavage of a C-terminal dipeptide of a substrate by cathepsin X can be observed under conditions that preclude efficient monopeptidyl carboxypeptidase activity. In addition, an inhibitor designed to exploit the unique structural features responsible for the carboxypeptidase activity of cathepsin X has been synthesized and tested against cathepsins X, B and L. Although of moderate potency, this E-64 derivative is the first reported example of a cathepsin X-specific inhibitor. By comparison, CA074 was found to inactivate cathepsin B at least 34 000-fold more efficiently than cathepsin
found to inactivate cathepsin B at least 34 000-fold more efficiently than cathepsin
DRUG DESCRIPTORS:
* cysteine proteinase--endogenous compound--ec; *cathepsin B --endogenous
compound--ec; *enzyme inhibitor--drug development--dv; *enzyme
inhibitor--pharmacology--pd
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec; ca 074--drug
development--dv; ca 074--pharmacology--pd
  4/3,K/38 (Item 37 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                   EMBASE No: 2001079403
11062658
Human cathepsin X: A novel cysteine protease with unique specificity
Menard R.; Nagler D.K.; Zhang R.; Tam W.; Sulea T.; Purisima E.O. R. Menard, Biotechnology Research Institute, National Research Council of Canada, 6100 Avenue Royalmount, Montreal, Que. H4P 2R2 Canada Advances in Experimental Medicine and Biology (ADV. EXP. MED. BIOL.) (United
                2000 , 477/- (317-322)
                    IŚSN: 0065-2598
CODEN: AEMBA
Document Type: Journal; Conference Paper
Language: ENGLISH
Number Of References: 14
Human cathepsin X: A novel cysteine protease with unique specificity
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec
  4/3,K/39 (Item 38 from file: 73) Links
     Fúlltext available through:
                                               USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                   EMBASE No: 2001079395
Review: Novel cysteine proteases of the papain family
Buhling F.; Fengler A.; Brandt W.; Welte T.; Ansorge S.; Nagler D.K.
F. Buhling, Institue of Immunology, Otto von Guericke Univ. Magdeburg, Magdeburg
Germany
Advances in Experimental Medicine and Biology (ADV. EXP. MED. BIOL.) (United
States ) 2000 , 477/- (241-254)
CODEN: AEMBA ISSN: 0065-2598
Document Type: Journal; Conference Paper
Language: ENGLISH
```

Page 58

```
Number Of References: 69
DRUG DESCRIPTORS:
* cysteine proteinase--endogenous compound--ec; *papain--endogenous compound--ec
cathepsin F--endogenous compound--ec; cathepsin K--endogenous compound--ec;
unclassified drug
Drug Terms (Uncontrolled): cathepsin o--endogenous compound--ec; cathepsin
V--endogenous compound--ec; cathepsin w--endogenous compound--ec; cathepsin
x--endogenous compound--ec
 4/3,K/40 (Item 39 from file: 73) Links
    Fulltext available through: USPTO Full Text Retrieval Options
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11023122
                 EMBASE No: 2000123291
Proteolytic signals from Magdeburg
Ansorge S.; Langner J.; Buhling F.; Lendeckel U.
S. Ansorge, Inst. of Experimental Internal Med., Otto-von-Guericke University,
D-39120 Magdeburg Germany
Immunology Today ( IMMUNOL. TODAY ) ( United Kingdom )
CODEN: IMTOD ISSN: 0167-5699
Publisher Item Identifier: S0167569900015863
Document Type: Journal; Article
Language: ENGLISH Summary Language: ENGLISH
                                                                          2000 , 21/4 (166-167)
DRUG DESCRIPTORS:
* microsomal aminopeptidase--endogenous compound--ec; *dipeptidyl peptidase
IV--endogenous compound--ec; *enzyme inhibitor--drug development--dv; *enzyme inhibitor--pharmacology--pd; *cathepsin --endogenous compound--ec
cathepsin S--endogenous compound--ec; cathepsin L--endogenous compound--ec;
cathepsin B--endogenous compound--ec; cathepsin H--endogenous compound--ec; cathepsin K--endogenous compound--ec; cathepsin D--endogenous compound--ec;
unclassified drug
Drug Terms (Uncontrolled): peptidase inhibitor--drug development--dv; peptidase
inhibitor --pharmacology--pd; cathepsin w--endogenous compound--ec; cathepsin F--endogenous compound--ec; cathepsin x--endogenous compound--ec
 4/3,K/41 (Item 40 from file: 73) Links
    Fulltext available through:
                                           USPTO Full Text Retrieval Options
EMBASE
(c) 2007 Elsevier B.V. All rights reserved.
                 EMBASE No: 2000429272
10939353
Flow cytometric analysis of enzymes in live spermatozoa before and after cryostorage
Schaller J.; Glander H.-J.
Dr. J. Schaller, Dermatohistological Unit, Department of Dermatology, St. Barbara
Hospital, Barbarastr. 67, 47167 Duisburg Germany Andrologia (ANDROLOGIA) (Germany) 2000, 32/0
                                                    <sup>2000</sup> , 32/6 (357-364)
                   ISSN: 0303-4569
CODEN: ANDRD
Document Type: Journal; Article
Language: ENGLISH Summa
Number Of References: 27
                          Summary Language: ENGLISH
...for butyryl esterase (P<0.05), prolyl-aminopeptidase (P<0.001) and val-lys-(VK)-cathepsin (P<0.001) most probably due to elevated enzyme activities.
The activities of FDA-esterase (P...
DRUG DESCRIPTORS:
* peptidase--endogenous compound--ec; *proteinase--endogenous compound--ec; *esterase--endogenous compound--ec; *collagenase--endogenous compound --ec; *collagenase--endogenous compound --ec
fluorescein; rhodamine 110; microsomal aminopeptidase--endogenous compound--ec;
                                                  Page 59
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cathepsinsearch.txt
subtilisin--endogenous compound--ec; dipeptidyl peptidase--endogenous compound--ec;
proline iminopeptidase --endogenous compound--ec; cathepsin--endogenous compound--ec
 4/3,K/42 (Item 41 from file: 73) Links
    Fulltext available through:
                                       USPTO Full Text Retrieval Options
(c) 2007 Elsevier B.V. All rights reserved.
10839518 EMBASE No: 2000320402
Characterization of TH1 and CTSZ, two non-imprinted genes downstream of GNAS1 in
chromosome 20q13
Bonthron D.T.; Hayward B.E.; Moran V.; Strain L. D.T. Bonthron, Molecular Medicine Unit, University of Leeds, St. James's University
Hospital, Leeds LS9 7TF
                              United Kingdom
Author Email: D.T.Bonthron@leeds.ac.uk
                                                     2000 , 107/2 (165-175)
Human Genetics ( HUM. GENET. ) ( Germany )
                 ISSN: 0340-6717
CODEN: HUGED
Document Type: Journal; Article
Language: ENGLISH
                       Summary Language: ENGLISH
Number Of References: 26
...probably not imprinted. Immediately downstream of TH1 lies CTSZ, encoding the
recently described cysteine protease, cathepsin Z. We have also elucidated the
genomic structure of this gene; it has six exons spanning...
DRUG DESCRIPTORS:
* cathepsin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin z--endogenous compound--ec
 4/3,K/43 (Item 42 from file: 73) Links
    Fulltext available through:
                                       USPTO Full Text Retrieval Options
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               EMBASE No: 2000316209
10834786
Biochemical characterization of human cathepsin X revealed that the enzyme is an
exopeptidase, acting as carboxymonopeptidase or carboxydipeptidase
Klemencic I.; Carmona A.K.; Cezari M.H.S.; Juliano M.A.; Juliano L.; Guncar G.; Turk
D.; Krizaj I.; Turk V.; Turk B.
B. Turk, Dept. of Biochemistry/Molec. Biol., Josef Stefan Institute, Jamova 39, 1000
              Slovenia
Ljubljana
Author Email: boris.turk@ijs.si
European Journal of Biochemistry ( EUR. J. BIOCHEM. ) ( United Kingdom )
                                                                                         2000
267/17 (5404-5412)
CODEN: EJBCA
                 ISSN: 0014-2956
Document Type: Journal; Article
Language: ENGLISH
                       Summary Language: ENGLISH
Number Of References: 44
Biochemical characterization of human cathepsin X revealed that the enzyme is an
exopeptidase, acting as carboxymonopeptidase or carboxydipeptidase
Cathepsin X, purified to homogeneity from human liver, is a single chain glycoprotein with a molecular mass of approx. eq. 33 kDa and pI 5.1-5.3. Cathepsin X was inhibited by stefin A, cystatin C and chicken cystatin (K(i) = 1.7-15... ...was also inhibited by two specific synthetic cathepsin B inhibitors, CA-074 and
GFG-semicarbazone. Cathepsin X was similar to cathepsin B and found to be a
carboxypeptidase with preference for a positively charged Arg in P1 position.
Contrary to the preference of cathepsin B, cathepsin X normally acts as a carboxymonopeptidase. However, the preference for Arg in the P1 position is so
```

strong that cathepsin X cleaves substrates with Arg in antepenultimate position, acting also as a carboxydipeptidase. A large hydrophobic......P1' position, although the enzyme cleaved all P1' residues investigated (Trp, Phe, Ala, Arg, Pro).

Page 60

cathepsinsearch.txt Cathepsin X also cleaved substrates with amide-blocked C-terminal carboxyl group with rates similar to those of the unblocked substrates. In contrast, no endopeptidase activity of cathepsin X could be detected on a series of o-aminobenzoic acid-peptidyl-N-[2,-dinitrophenyl]ethylenediamine... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *carboxypeptidase--endogenous compound--ec; *dipeptidase--endogenous compound--ec liver enzyme--endogenous compound--ec; stefin A; cystatin C; stefin B; kininogen; semicarbazone; cathepsin B; tryptophan; phenylalanine; arginine; proline... 4/3,K/44 (Item 43 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000138039 10728641 Role for cathepsin F in invariant chain processing and major histocompatibility complex class II peptide loading by macrophages Shi G.-P.; Bryant R.A.R.; Riese R.; Verhelst S.; Driessen C.; Li Z.; Bromme D.: Ploegh H.L.; Chapman H.A. H.A. Chapman, Pulmonary and Critical Care Div., University of California, 505 Parnassus Ave., San Francisco, CA 94143-0111 United States Author Email: halchap@itsa.ucsf.edu Journal of Experimental Medicine (J. EXP. MED.) (United States) 03 APR 2000 . 191/7 (1177-1185) CODEN: JEMEA IS ISSN: 0022-1007 Document Type: Journal ; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 42 ...and dendritic cells revealed two enzymes expressed exclusively in macrophages, cathepsins Z and F. Recombinant cathepsin Z did not generate CLIP from Ii-MHC class II complexes, whereas cathepsin F was as... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *major histocompatibility antigen class 2--endogenous compound--ec; *cell membrane protein --endogenous compound--ec cysteine proteinase--endogenous compound--ec; recombinant enzyme; cathepsin S; unclassified drug Drug Terms (Uncontrolled): cathepsin F--endogenous compound--ec; class ii associated invariant chain peptide--endogenous compound--ec; cathepsin z 4/3,K/45 (Item 44 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000192563 The new subfamily of cathepsin-Z-like protease genes includes Tc-cpz-1, a cysteine protease gene expressed in Toxocara can's adults and infective stage larvae Falcone F.H.; Tetteh K.K.A.; Hunt P.; Blaxter M.L.; Loukas A.; Maizels R.M. R.M. Maizels, Inst. Cell Animal/Population Biol., University of Edinburgh, West Mains Road, Edinburgh EH9 3JT (Author Email: r.maizels@ed.ac.uk United Kinadom Experimental Parasitology (EXP. PARASITOL.) (United States) 2000 , 94/3 (201-207)ISSN: 0014-4894 CODEN: EXPAA Document Type: Journal : Article Language: ENGLISH Number Of References: 32 The new subfamily of cathepsin-Z-like protease genes includes Tc-cpz-1, a cysteine protease gene expressed in Toxocara canis...

Page 61

DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin tc cpz 1--endogenous compound--ec 4/3,K/46 (Item 45 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000116176 Mouse cathepsin M, a placenta-specific lysosomal cysteine protease related to cathepsins L and P Sol-Church K.; Frenck J.; Mason R.W.
R.W. Mason, Laboratory of Enzymology, Department of Research, Alfred I.du Pont Hosp.
for Children, P.O. Box 269, Wilmington, DE 19899 United States
Author Email: rmason@nemours.org Biochimica et Biophysica Acta - Gene Structure and Expression (BIOCHIM. BIOPHYS. ACTA GENE STRUCT. EXPR.) (Netherlands) CODEN: BBGSD ISSN: 0167-4781 25 APR 2000 , 1491/1-3 (289-294) Publisher Item Identifier: S0167478100000300 Document Type: Journal ; Article Language: ENGLISH Summa Number Of References: 34 Summary Language: ENGLISH DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *complementary DNA--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec cathepsin L--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin m--endogenous compound--ec; cathepsin p --endogenous compound--ec 4/3,K/47 (Item 46 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 2000116156 10651103 Murine and human cathepsin Z: cDNA-cloning, characterization of the genes and chromosomal localization Deussing J.; Von Olshausen I.; Peters C. C. Peters, Institut Molekular Medizin, Klinikum, Albert-Ludwig-Universitat, Hugstetter Strasse 55, 79106 Freiburg Germany Hugstetter Strasse 33, 73100 Fielding Author Email: peters@mm11.ukl.uni-freiburg.de
Biochimica et Biophysica Acta - Gene Structure and Expression (BIOCHIM. BIOPHYS.
ACTA GENE STRUCT. EXPR.) (Netherlands) 25 APR 2000 , 1491/1-3 (93-106) ACTA GENE STRUCT. EXPR.) (Netherlands) CODEN: BBGSD ISSN: 0167-4781 Publisher Item Identifier: S016747810000021X Document Type: Journal ; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 53 Murine and human cathepsin Z: cDNA-cloning, characterization of the genes and chromosomal localization ...encoding a predicted polypeptide of 306 amino acids was characterized. The new protease, tentatively named cathepsin Z, exhibits all features characteristics of a papain-like cysteine protease, including the highly conserved residues of the 'catalytic triad'. Cathepsin Z shares only 26-35% overall homology with previously described mammalian papain-like cysteine peptidases and... ...within the family of papain-like cysteine peptidases. Genomic clones covering the murine and human cathepsin Z genes were isolated. They comprise six exons and five introns spanning a 12-kb region of genomic DNA, respectively. Murine cathepsin Z was mapped to chromosome 2, a region with synteny homology to a region of human chromosome 20 to

Page 62

which human cathepsin Z has been mapped previously. Northern blot analysis revealed ubiquitous expression of murine cathepsin Z. Multiple transcriptional start sites were identified for the murine cathepsin Z gene and together with the absence of a TATA box, a high G+C content.....CpG island and the presence of several sp1-binding sites in the promoter region, murine cathepsin Z may be classified as a 'housekeeping' gene. Copyright (C) 2000 Elsevier Science B.V. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec complementary DNA--endogenous compound--ec

4/3,K/48 (Item 47 from file: 73) Links

Fulltext available through: USPTO Full Text Retrieval Options

EMBASE

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10633942 EMBASE No: 2000099482

Crystal structure of cathepsin X: A flip-flop of the ring of His23 allows carboxy-monopeptidase and carboxy-dipeptidase activity of the protease

Guncar G.; Klemencic I.; Turk B.; Turk V.; Karaoglanovic-Carmona A.; Juliano L.; Turk D.

D. Turk, Dept. of Biochem./Molecular Biology, Jozef Stefan Institute, Jamova 39, 1000 Ljubljana Slovenia

Author Email: dusan.turk@ijs.si

Structure (STRUCTURE) (United Kingdom) 15 MAR 2000, 8/3 (305-313)

CODEN: STRUE ISSN: 0969-2126 Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 54

Crystal structure of cathepsin X: A flip-flop of the ring of His23 allows carboxy-monopeptidase and carboxy-dipeptidase activity...

Background: Cathepsin X is a widespread, abundantly expressed papain-like mammalian lysosomal cysteine protease. It exhibits carboxy-monopeptidase.....of the two enzyme activities has actually been monitored. Results: The crystal structure of human cathepsin X has been determined at 2.67 Angstrom resolution. The structure shares the common features of....like enzyme fold, but with a unique active site. The most pronounced feature of the cathepsin X structure is the mini-loop that includes a short three-residue insertion protruding into the.....terminal carboxyl group of a substrate in two different sidechain conformations. Conclusions: The structure of cathepsin X exhibits a binding surface that will assist in the design of specific inhibitors of cathepsin X as well as of cathepsin B and thereby help to clarify the physiological roles of...

DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *histidine--endogenous compound--ec; *peptidase--endogenous compound--ec; * proteinase--endogenous compound--ec

4/3,K/49 (Item 48 from file: 73) Links

Fulltext available through: USPTO Full Text Retrieval Options

EMBASE

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10601951 EMBASE No: 2000067209

Cathepsin Q, a novel lysosomal cysteine protease highly expressed in placenta

Sol-Church K.; Frenck J.; Mason R.W.

R.W. Mason, Láboratory of Enzymology, Department of Research, Alfred I. duPont Hospital Children, PO Box 269, Wilmington, DE 19899 United States

Author Email: rmason@nemours.org

Biochemical and Biophysical Research Communications (BIOCHEM. BIOPHYS. RES. COMMUN.) (United States) 27 JAN 2000, 267/3 (791-795)

CODEN: BBRCA ISSN: 0006-291X

Document Type: Journal; Article

cathensinsearch.txt Summary Language: ENGLISH Language: ENGLISH Number Of References: 22 ...is predicted that cathepsin Q will differ in catalytic specificity to another placental-specific protease, cathepsin P, indicating that these enzymes will have unique proteolytic functions in extra-embryonic tissues. (C) 2000... DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec; *lysosome enzyme--endogenous compound--ec 4/3,K/50 (Item 49 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1999415968 10531360 Cathepsin Y (a novel thiol enzyme) produces kinin potentiating peptide from the component protein of rat plasma Sakamoto E.; Sakao Y.; Taniguchi Y.; Yamafuji K. E. Sakamoto, Department of Food and Nutrition, Nakamura Gakuen University, Jonan-ku, Fukuoka 814-0198 Japan Immunopharmacology (IMMUNOPHARMACOLOGY) (Netherlands)
CODEN: IMMUD ISSN: 0162-3109
Publisher Item Identifier: S016231099900079X 1999 , 45/1-3 (207-214) Document Type: Journal; Article Language: ENGLISH Summary Language: ENGLISH Number Of References: 18 Cathepsin Y (a novel thiol enzyme) produces kinin potentiating peptide from the component protein of rat plasma Rat spleen cathepsin Y (a novel enzyme) that produces bradykinin (BK) potentiating peptide (BPP) from rat plasma was isolated....from cDNA cloned by reverse transcription-polymerase chain reaction (RT-PCR). We propose the name cathepsin Y for this enzyme considering its origin, characteristics and the amino acid sequence. BPP potentiates not....when the level is doubled. The precursor proteins that produce BPP by the action of cathepsin Y are eluted into two fractions when the heated plasma was applied to a negative ion....this paper, we report on the characteristics and the amino acid sequence of rat spleen cathepsin Y, its structure and the potentiating activity of BPP, and isolation of the precursor protein. Copyright.. DRUG DESCRIPTORS: * cathepsin--endogenous compound--ec; *bradykinin--endogenous compound--ec; *thiol proteinase--endogenous compound--ec; * kinin--endogenous compound--ec
Drug Terms (Uncontrolled): cathepsin y--endogenous compound--ec 4/3,K/51 (Item 50 from file: 73) Links USPTO Full Text Retrieval Options Fulltext available through: **EMBASE** (c) 2007 Elsevier B.V. All rights reserved. EMBASE_No: 1999379729 07906071 Cathepsin P, a novel protease in mouse placenta Sol-Church K.; Frenck J.; Troeber D.; Mason R.W. R.W. Mason, Laboratory of Enzymology, Department of Research, Alfred I. duPont Hospital Children, PO Box 269, Wilmington, DE 19899 United States Author Email: rmason@nemours.org Biochemical Journal (BIOCHEM. J.) (United Kingdom) 15 OCT 1999 . 343/2 (307 - 309)

Page 64

CODEN: BÍJOA ISSN: 0264-6021 Document Type: Journal ; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 13 Cathepsin P, a novel protease in mouse placenta

The complete cDNA nucleotide sequence of a novel cathepsin derived from mouse placenta, termed cathepsin P, was determined. mRNA for cathepsin P was expressed in placenta and at lower levels in visceral yolk sac, but could not... DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec proteinase--endogenous compound--ec; complementary DNA--endogenous compound--ec; messenger RNA--endogenous compound--ec

4/3,K/52 (Item 51 from file: 73) Links Fúlltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1999344211

Human cathepsin X: A cysteine protease with unique carboxypeptidase activity

Nagler D.K.; Zhang R.; Tam W.; Sulea T.; Purisima E.O.; Menard R. R. Menard, Biotechnology Research Institute, National Research Council of Canada, 6100 Royalmount Ave., Montreal, Que. H4P 2R2 Canada Author Email: robert.menard@nrc.ca

Biochemistry (BIOCHEMISTRY) (United States) 28 SEP 1999 , 38/39 (12648-12654)

ISSN: 0006-2960 CODEN: BICHA Document Type: Journal ; Article

Language: ENGLISH Summary Language: ENGLISH

Number Of References: 40

Human cathepsin X: A cysteine protease with unique carboxypeptidase activity

Cathepsin X is a novel cysteine protease which was identified recently from the EST (expressed sequence tags) database. In a homology model of the mature cathepsin X, a unique three residue insertion between the Gln22 of the oxyanion hole and the active....verify this hypothesis, human procathepsin X was expressed in Pichia pastoris and converted to mature cathepsin X using small amounts of human cathepsin L. Cathepsin X was found to display excellent carboxypeptidase activity against the substrate Abz-FRF(4NOinf 2), with.....1 ssup -sup 1 at the optimal pH of 5.0. However, the activity of cathepsin X against the substrates Cbz-FR-MCA and Abz-AFRSAAQ-EDDnp was found to be extremely....k(cat)/K(M) values lower than 70 Msup -sup 1 ssup -sup 1. Therefore, cathepsin X displays a stricter exopeptidase activity than cathepsin B. No inhibition of cathepsin X by cystatin C could be detected up to a concentration of 4 muM of inhibitor... ... the bound carboxypeptidase substrate is predicted to establish a number of favorable contacts within the cathepsin X binding site, in particular with residues His23 and Tyr27 from the mini-loop. The presence substrates in the primed subsites of the protease. The marked structural and functional differences of cathepsin X relative to other members of the papain family of cysteine proteases will be of great... DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec; *cysteine proteinase--endogenous compound--ec; *carboxypeptidase--endogenous compound--ec; * cathepsin l--endogenous compound--ec; *cathepsin b--endogenous compound--ec histidine--endogenous compound--ec; tyrosine--endogenous compound--ec; cystatin c--endogenous compound--ec; cysteine
Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/53 (Item 52 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1998297006 07403528 Human cathepsin X: A novel cysteine protease of the papain family with a very short

proregion and unique insertions

Nagler D.K.; Menard R. R. Menard, Biotechnology Research Institute, National Research Council Canada, 6100 Avenue Royalmount, Montreal, Que. H4P 2R2 Canada FEBS Letters (FEBS LETT.) (Netherlands) 1998 CODEN: FEBLA ISSN: 0014-5793 1998 , 434/1-2 (135-139) Publisher Item Identifier: S0014579398009648 Document Type: Journal ; Article Summary Language: ENGLISH Language: ENGLISH Number of References: 40 Human cathepsin X: A novel cysteine protease of the papain family with a very short proregion and unique... A novel cDNA encoding a cysteine protease of the papain family named cathepsin X was obtained by PCR amplification from a human ovary cDNA library. The cathepsin X cDNA is ubiquitously expressed in human tissues and contains an open reading frame of 912... ...highly conserved regions in papain-like cysteine proteases including the catalytic residues are present in cathepsin X. The mature part of cathepsin X is 26-32% identical to human cathepsins B, C, H, K, L, O, S and W. The cathepsin X sequence contains several unique features: (i) a very short proregion; (ii) a three amino acid... DRUG DESCRIPTORS: cathepsin: *papain--endogenous compound--ec; *cysteine proteinase --endogenous compound--ec cathepsin s--endogenous compound--ec; cathepsin b--endogenous compound--ec 4/3,K/54 (Item 53 from file: 73) Links Fulltext available through: USPTO Full Text Retrieval Options (c) 2007 Elsevier B.V. All rights reserved. EMBASE No: 1998227542 Cathepsin Z, a novel human cysteine proteinase with a short propeptide domain and a unique chromosomal location Santamaria I.; Velasco G.; Pendas A.M.; Fueyo A.; Lopez-Otin C. C. Lopez-Otin, Depto. de Bioquimica/Biólogia Molec., Facultad de Medicina, Universidad de Oviedo, 33006 Oviedo Spain Author Email: CLO@DWARF1.QUIMICA.UNIOVI.ES Journal of Biological Chemistry (J. BIOL. CHEM.) (United States) 03 JUL 1998, 273/27 (16816-16823) CODEN: JBCHA ISSN: 0021-9258 Document Type: Journal; Article Language: ENGLISH Summ Number Of References: 62 Summary Language: ENGLISH Cathepsin Z, a novel human cysteine proteinase with a short propeptide domain and a unique chromosomal location ...revealed that the isolated cDNA codes for a polypeptide of 303 amino acids, tentatively called cathepsin Z, that exhibits structural features characteristic of cysteine proteinases. Fluorescent in situ hybridization experiments revealed that the human cathepsin Z gene maps to chromosome 20q13, a location that differs from all cysteine proteinase genes mapped to date. The cDNA encoding cathepsin Z was expressed in Escherichia coli as a fusion protein with glutathione S- transferase, and after... ...amido-4- methylcoumarin, used as a substrate for cysteine proteinases. Northern blot analysis demonstrated that cathepsin Z is widely expressed in human tissues, suggesting that this enzyme could be involved in the normal intracellular protein degradation taking place in all cell types. Cathepsin Z is also ubiquitously distributed in cancer cell lines and in primary tumors from different sources.

chromosomal location among cysteine proteinases, we propose that cathepsin Z may be the first representative of a novel subfamily of this class of proteolytic enzymes.

different sources... ...unusual short propeptide, together with its unique

DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec Drug Terms (Uncontrolled): cathepsin z--endogenous compound--ec

4/3,K/55 (Item 54 from file: 73) Links
Fulltext available through: USPTO Full Text Retrieval Options
EMBASE
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06200066 EMBASE No: 1995231005
Cloning and complete coding sequence of a povel human cathensin express

Cloning and complete coding sequence of a novel human cathepsin expressed in giant cells of osteoclastomas

Li Y.-P.; Alexander M.; Wucherpfennig A.L.; Yelick P.; Chen W.; Stashenko P. Forsyth Dental Center, 140 Fenway, Boston, MA 02115 United States Journal of Bone and Mineral Research (J. BONE MINER. RES.) (United States) 1995 , 10/8 (1197-1202)

CODEN: JBMRE ISSN: 0884-0431 Document Type: Journal; Article

Language: ENGLISH Summary Language: ENGLISH

...has been identified by differential screening of a human osteoclastoma cDNA library. This molecule, termed cathepsin X, appears to represent the human homolog of the osteoclast-expressed rabbit cathepsin OC-2. Cathepsin X (GenBank accession number U20280) is 93.9% identical to OC-2 at the amino acid level, and is 92% identical at the nucleotide level within the coding region. Cathepsin X is 52.2 and 46.9% identical to cathepsins S and L, respectively, and is therefore clearly distinct from these enzymes. Cathepsin X mRNA was localized to multinucleated giant cells within the osteoclastoma tumor by in situ hybridization. These data strongly support the hypothesis that cathepsin X represents a novel cysteine proteinase which is expressed at high levels in osteoclasts. DRUG DESCRIPTORS:

* cathepsin--endogenous compound--ec cysteine proteinase--endogenous compound--ec; unclassified drug Drug Terms (Uncontrolled): cathepsin x--endogenous compound--ec

4/3,K/56 (Item 1 from file: 35) Links
Dissertation Abs Online
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01980017 ORDER NO: AADAA-IMQ83852
The design of substrates for cathepsin X

Author: Devanathan, Gopal Degree: M.Sc.

Degree: M.Sc. Year: 2003

Corporate Source/Institution: Concordia University (Canada) (0228)

Source: Volume 42/03 of MASTERS ABSTRACTS. of Dissertations Abstracts International.

PAGE 941 . 90 PAGES ISBN: 0-612-83852-8

The design of substrates for cathepsin X

...diseases such as arthritis, Alzheimer's, and cancer, they are attractive targets for inhibitor design. Cathepsin X is a cysteine protease that was only recently discovered. The primary structure of cathepsin X contains several unique features that clearly distinguish it from the other human cysteine proteases. The... ... a systematic study on the S2, S1, and S1<super>′</super> subsites of the cathepsin X active site and to gain a detailed understanding of the enzyme's substrate specificity.

Three libraries of compounds have been synthesized based on the parent compound 2-Abz-Phe-Arg-Phe(4NO₂). In each library, the 20 natural... ...prime;</super> sites respectively, while keeping the other positions fixed. In reference to the parent compound, P2 is occupied by Phe, P1 by Arg, and Page 67

P1<super>′</super> by Phe.....by docking 2-Abz-Phe-Arg-Phe(4NO₂) and analogues to the cathepsin X active site in order to gain a detailed understanding of factors underlying substrate specificity. Knowledge...

4/3,K/57 (Item 1 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0377298 DBA Accession No.: 2005-23004 PATENT Identification of a compound capable of modulating the activity of cathepsin Z in a cell comprises measuring the cell's base level of cathepsin Z activity in the absence and presence of the compound involving vector-mediated gene transfer and expression in host cell for therapy Author: SALTZMAN A G; TANG Z; PALEJWALA V; CAVALLO J Patent Assignee: AVENTIS PHARM INC 2005 Patent Number: WO 200565693 Patent Date: 20050721 WPI Accession No.: 2005-533570 (200554) Priority Application Number: US 533330 Application Date: 20031230 National Application Number: WO 2004US41815 Application Date: 20041214 Language: English Identification of a compound capable of modulating the activity of cathepsin Z in a cell comprises measuring the cell's base level of cathepsin Z activity in the absence and presence of the compound involving vector-mediated gene transfer and expression in host cell for therapy Abstract: DERWENT ABSTRACT: NOVELTY - Identification of a compound (I) capable of modulating the activity of cathepsin Z in a cell comprises measuring the cell's base level of cathepsin Z activity in the absence of a candidate compound; introducing the candidate compound; and measuring the cell's level of cathepsin Z activity in the presence of the candidate compound. DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for: (1) the compound (I) capable of modulating activity of cathepsin Z; and (2) a pharmaceutical comprising (I) and excipient for treating an inflammatory. ACTIVITY - Antiinflammatory; Immunosuppressive; Antiarthritic; Antirheumatic; Neuroprotective. MECHANISM OF ACTION - Cathepsin Z modulator. Test details are described but no results given. USE - (I) is useful to treat... Descriptors: recombinant cathepsin-Z prep., isol., vector-mediated gene transfer, expression in host cell, appl., inflammatory disease, autoimmune disease...

4/3,K/58 (Item 2 from file: 357) Links
Derwent Biotech Res.
(c) 2007 The Thomson Corp. All rights reserved.
0338230 DBA Accession No.: 2004-10522 PATENT
Use of polynucleotide sequence encoding Cathespin Y protein for identification of therapeutic agent useful for treating stroke e.g. ischemic stroke vector-mediated cathespin-Y gene transfer, expression in host cell and antisense oligonucleotide for drug screening and gene therapy

Author: LUBBERT H; ZWILLING S; ENGELS P

Patent Assignee: LUBBERT H; ZWILLING S; ENGELS P 2003

Patent Number: US 20030232740 Patent Date: 20031218 WPI Accession No.: 2004-142033 (200414)

Priority Application Number: US 392809 Application Date: 20030319 National Application Number: US 392809 Application Date: 20030319 Language: English

Abstract: ...of potential therapeutic agent for treating stroke involves contacting a cell capable of expressing a Cathepsin Y gene or homologues or fragments with the potential therapeutic agent; detecting a level of expression of the Cathepsin Y gene in the test cell; comparing expression in the test cell to a reference cell.....of potential therapeutic agent for treating stroke involves contacting a cell capable of expressing a Cathepsin Y gene or homologues or fragments with the potential therapeutic agent; detecting a level of expression of the Cathepsin Y gene in the test cell; comparing the level of expression of the Cathepsin Y gene in the test

cathepsinsearch.txt

cell to a level of expression of the Cathepsin Y gene in a reference cell whose disease stage is known; and identifying the difference in the expression level of the Cathepsin Y gene in the test cell and the reference cell. INDEPENDENT CLAIMS are included for the following: (a) a composition comprising a compound of formula (I) or its salt; (b) a composition comprising a nucleic acid sequence (S1) which is an antisoned sequence (S2) encoding Cathepsin Y antisense sequence compared to a nucleic acid sequence (S2) encoding Cathepsin Y, its homologue or fragment. (S2) Has sequence of 1140 or 1500 nucleotide bases as given....is 0, then R4 is other than -N(CH3)OCH3. ACTIVITY - Cerebroprotective. MECHANISM OF ACTION - Cathepsin Y protein inhibitor. Test details are described, but no results are given. USE - For identifying potential...

4/3,K/59 (Item 3 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0335363 DBA Accession No.: 2004-07655 PATENT New antisense compound targeted to nucleic acid molecules encoding cathepsin Z, useful for treating diseases associated with expression of cathepsin Z, e.g. encephalitis, viral infection, or hyperproliferative disorder involving vector-mediated gene transfer and expression in host cell for use in therapy Author: DOBIE K W Patent Assignee: ISIS PHARM INC 2003
Patent Number: US 20030224511 Patent Date: 20031204 WPI Accession No.: 2004-060543 (200406)Priority Application Number: US 159266 Application Date: 20020531 National Application Number: US 159266 Application Date: 20020531 Language: English New antisense compound targeted to nucleic acid molecules encoding cathepsin Z. useful for treating diseases associated with expression of cathepsin Z, e.g. encephalitis, viral infection, or hyperproliferative disorder involving vector-mediated gene transfer and expression... Abstract: DERWENT ABSTRACT: NOVELTY - A compound (I) 8-80 nucleobases in length targeted to a nucleic acid molecule encoding cathepsin Z, is new. DETAILED DESCRIPTION - A compound (I) 8-80 nucleobases in length targeted to a nucleic acid molecule encoding cathepsin Z, is new. The compound specifically hybridizes with the nucleic acid molecule encoding cathepsin Z and inhibits the expression of cathepsin Z, or specifically hybridizes with at least an 8-nucleobase portion of a preferred target region on a nucleic acid molecule encoding cathepsin Z. INDEPENDENT CLAIMS are included for the following: (1) a composition comprising (I) and a pharmaceutical carrier or diluent; (2) a method of inhibiting the expression of cathepsin Z in cells or tissues comprising contacting the cells or tissues with (I); and (3) a method of treating an animal having a disease or condition associated with cathepsin Z comprising administering to the animal a therapeutic or prophylactic amount of (I) so that expression of cathepsin Z is inhibited. BIOTECHNOLOGY -Preparation: The antisense compounds are produced by solid phase synthesis Preferred Compound: The compound is an antisense oligonucleotide, preferably a chimeric oligonucleotide. The antisense oligonucleotide comprises: (a) at least... .. Inhibitor Z. USE - The antisense oligonucleotides and compounds are useful for inhibiting the expression of cathepsin Z, and for treating diseases or conditions associated with expression of cathepsin Z, e.g. encephalitis, viral infection, or hyperproliferative disorder, such as cancer (all claimed). The antisense... Descriptors: recombinant cathepsin-Z prep., isol., vector-mediated gene transfer, expression in host cell, antisense oligonucleotide, appl. encephalitis, virus...

4/3,K/60 (Item 4 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0334375 DBA Accession No.: 2004-06667 PATENT Composition useful for treating pain e.g. neuropathic pain comprises polynucleotide sequence sense and antisense sequence for use in disease therapy and gene therapy

Author: LUBBERT H; ENGELS P; SCHMITZ B
Patent Assignee: LUBBERT H; ENGELS P; SCHMITZ B 2003
Patent Number: US 20030212003 Patent Date: 20031113 WPI Accession No.: 2004-041675

(200404)

Priority Application Number: US 369386 Application Date: 20030214 National Application Number: US 369386 Application Date: 20030214

Language: English

...therapeutic agents for treating pain involving: (a) contacting a test cell capable of expressing a Cathepsin Y gene, its homologues or fragments with the potential therapeutic agent; (b) detecting a level of expression of the Cathepsin Y gene in the test cell; (c) comparing the level of expression of the Cathepsin Y gene in the test cell to that in a reference cell; and (d) identifying the difference in the expression levels of the Cathepsin Y gene in the test cell and reference cell; (2) identification of a therapeutic agent for treating pain involving: (a) incubating a sample comprising a Cathepsin Y protein, a test compound/agent and a polymentide which is a target of Cathepsin Y protein proteolysis: (h) determining an polypeptide which is a target of Cathepsin Y protein proteolysis; (b) determining an aminoterminal amino acid of a peptide resulting from the proteolysis... ...nucleic acid sequence is an antisense sequence compared to a nucleic acid sequence that encodes Cathepsin Y and has a sequence of 1387 or 1500 nucleotide bases. BIOTECHNOLOGY - Preferred Method: The expression of the Cathepsin Y gene is determined by at least one method selected from PCR of a cDNA, hybridizing a sample DNA and detecting a Cathepsin Y protein. ACTIVITY - Analgesic; Antidiabetic; Neuroprotective; Virucide; Vulnerary; Cardiant. MECHANISM OF ACTION - Cathepsin Y protein inhibitor. No biological data given. USE - The compound is useful for treating pain e.g. neuropathic pain (claimed), diabetic neuropathy, post-herpetic neuralgia....reflex sympathetic dystrophy and causalgia, myocardial syndromes or idiopathic pain. ADVANTAGE - The composition efficiently downregulates Cathepsin Y activity and hence treats pain. EXAMPLE - No relevant example given. (22 pages) Descriptors: polynucleotide composition, cathepsin Y gene, antisense sequence, polymerase chain reaction, appl. pain, neuropathic pain, diabetic neuropathy, post-herpetic neuralgia...

4/3,K/61 (Item 5 from file: 357) Links

Derwent Biotech Res.

(c) 2007 The Thomson Corp. All rights reserved. 0328457 DBA Accession No.: 2004-00749 PATENT

Developing medicament used for treating pain comprises using polynucleotide sequence encoding cathepsin Y involving vector-mediated gene transfer and expression in host cell for use in neuropathic pain therapy

Author: LUEBBERT H; SCHMITZ B

Patent Assignee: BIOFRONTERA PHARM AG 2003

Patent Number: EP 1336847 Patent Date: 20030820 WPI Accession No.: 2003-814978 (

2003)

Priority Application Number: EP 20023400 Application Date: 20020214 National Application Number: EP 20023400 Application Date: 20020214

Language: English

Developing medicament used for treating pain comprises using polynucleotide sequence encoding cathepsin Y involving vector-mediated gene transfer and expression in host

cell for use in neuropathic pain...

Abstract: DERWENT ABSTRACT: NOVELTY - Developing a medicament for treating pain comprises using a polynucleotide sequence encoding cathepsin Y. DETAILED DESCRIPTION - Developing a medicament for treating pain, for diagnosing pain status outside of a....efficacy of pain treatment outside of a living body, comprises using a polynucleotide sequence encoding cathepsin Y or homologs or fragments or the corresponding protein or peptide. INDEPENDENT CLAIMS are also included for: (1) the use of a compound downregulating cathepsin Y expression or activity for manufacture of a medicament for treatment of pain; (2) a diagnostic....isolated nucleic acid sequence comprising an 'antisense' sequence compared to a nucleic acid sequence encoding cathepsin Y or a fragment of the nucleic acid sequence as a medicament; and (4) a transgenic animal where the gene encoding cathepsin Y is manipulated in the animal in comparison to the wild type. ACTIVITY - Analgesic. MECHANISM OF ACTION -

Cathepsin Y inhibitor. Tests are described, but no results are given. USE - Used for treating pain, particularly...

Descriptors: recombinant cathepsin-Y prep., isol., vector-mediated gene transfer, expression in host cell, polymerase chain reaction, appl. neuropathic...

4/3,K/62 (Item 6 from file: 357) Links
Derwent Biotech Res.
(c) 2007 The Thomson Corp. All rights reserved.
0313249 DBA Accession No.: 2003-14389 PATENT
New transgenic mice comprising a disruption in a cathepsin Z (CTSZ) useful as models for diseases or conditions associated with phenotypes relating to a disruption in a CTSZ gene, and in identifying drugs for treating a disease vector-mediated mutant gene transfer and expression in embryonic stem cell for transgenic mouse construction for use as an animal model in disease therapy

Author: WISOTZKEY R G; KIRK C J Patent Assignee: DELTAGEN INC 2003

Patent Number: wo 200326403 Patent Date: 20030403 WPI Accession No.: 2003-354621

(200333)

Priority Application Number: US 324639 Application Date: 20010924 National Application Number: WO 2002US30506 Application Date: 20020924

Language: English
New transgenic mice comprising a disruption in a cathepsin Z (CTSZ) useful as models for diseases or conditions associated with phenotypes relating to a disruption...
Abstract: DERWENT ABSTRACT: NOVELTY - A transgenic mouse comprising a disruption in a cathepsin Z (CTSZ) gene, where there is no native expression of endogenous CTSZ gene, is new. DETAILED.....a pharmaceutical composition for a condition associated with a function of CTSZ, comprises identifying a compound that modulates CTSZ, synthesizing the identified compound, and incorporating the compound into a pharmaceutical carrier. USE - The transgenic mouse is useful as a model for diseases.....symptoms; and in testing and developing new treatments relating to behavioral phenotypes. EXAMPLE - Disruptions in cathepsin Z (CTSZ) genes were produced by homologous recombination. Transgenic mice comprising disruptions in CTSZ genes were...

Descriptors: transgenic mouse construction, vector-mediated mutant cathepsin- Z gene transfer, expression in embryonic stem cell, phenotyping, animal model, antagonist, agonist, database, homologous recombination...

4/3,K/63 (Item 7 from file: 357) Links
Derwent Biotech Res.
(c) 2007 The Thomson Corp. All rights reserved.
0257611 DBA Accession No.: 2000-12101 PATENT
New human cathepsin-Y protein, a gene encoding it and its application - diagnosis, therapy, gene therapy and drug screening

Corporate Source: Japan.

Patent Assignee: Fuji-Pharm. 2000

Patent Number: JP 2000157263 Patent Date: 20000613 WPI Accession No.: 2000-468198

(2041)

Priority Application Number: JP 98352110 Application Date: 19981126 National Application Number: JP 98352110 Application Date: 19981126

Language: Japanese

New human cathepsin-Y protein, a gene encoding it and its application Abstract: A human-derived cathepsin-Y protein (I) or a new human-derived cathepsin-Y protein which has at least 49% homology to the protein sequence of (I) has a... ... I) or its salts, peptides, etc., the DNA or the antibody; a drug containing a compound promoting or inhibiting biological activity of one of the claimed proteins, their partial peptides or...

Descriptors: human recombinant cathepsin-Y prep., cysteine protease act., monoclonal antibody, vector expression in host cell, DNA probe hybridization, appl...

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 6/3,K/1 (Item 1 from file: 98) Links
General Sci Abs
(c) 2007 The HW Wilson Co. All rights reserved.
            H.w. Wilson Record Number: BGSI98037731
03787731
Advances in Health Sciences Education: Theory and Practice.
Augmented Title: review of triannual journal edited by Henk Schmidt et al., Kluwer
Academic Publishers
Saltzman, Alan R
Byrd, Gary D
JAMA ( JAMA ) v. 279 no13 (Apr. 1 '98) p. 1045
ISSN: 0098-7484
Language: English
Country Of Publication: United States
Saltzman, Alan R
 6/3,K/2 (Item 1 from file: 143) Links
Biol. & Agric. Index
(c) 2007 The HW Wilson Co. All rights reserved.
            H.W. Wilson Record Number: BBAI95034200
                                           Page 72
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cathepsinsearch.txt Influence of cardiac action on gas mixing in closed-chest dogs

Zhang, Shaoping
Saltzman, Alan R; Klocke, Robert A
Journal of Applied Physiology v. 79 (July '95) p. 113-20
Document Type: Feature Article ISSN: 8750-7587
Saltzman, Alan R...

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 9/3,K/1 (Item 1 from file: 399) Links
CA SEARCH(R)
(c) 2007 American Chemical Society. All rights reserved.
                     CA: 143(7)109795k
143109795
                                                       PATENT
Cathepsin Z inhibitors for treatment of rheumatoid arthritis and other autoimmune
diseases
Inventor (Author): Saltzman, Alan G.; Tang, Zhihua; Palejwala, Vaseem; Cavallo, Jean
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Location: USA
Assignee: Aventis Pharmaceuticals Inc.
Patent: PCT International; WO 200565693 A2
                                                            Date: 20050721
Application: wo 2004us41815 (20041214) *US 2003PV533330 (20031230)
Pages: 28 pp.
CODEN: PIXXD2
Language: English
Patent Classifications:
  class:
             A61K-031/70A
Designated Countries: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY; BZ; CA;
CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; MA; MD; MG; MK; MN; MW; MX; MZ; NA; NI; NO; NZ; OM; PG; PH; PL; PT; RO; RU; SC; SD; SE; SG; SK;
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CA SEARCH(R)
(c) 2007 American Chemical Society. All rights reserved.
                       CA: 143(7)109795k
                                                           PATENT
Cathepsin Z inhibitors for treatment of rheumatoid arthritis and other autoimmune.
diseases
Inventor (Author): Saltzman, Alan G.; Tang, Zhihua; Palejwala, Vaseem; Cavallo, Jean
Location: USA
Assignee: Aventis Pharmaceuticals Inc.
Patent: PCT International; WO 200565693 A2 Date: 20050721 Application: WO 2004US41815 (20041214) *US 2003PV533330 (20031230)
Pages: 28 pp.
CODEN: PIXXD2
Language: English
Patent Classifications:
               A61K-031/70A
   Class:
Designated Countries: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY; BZ; CA;
CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; MA; MD; MG; MK; MN; MX; MZ; NA; NI; NO; NZ; OM; PG; PH; PL; PT; RO; RU; SC; SD; SE; SG; SK; SL; SY; TJ; TM; TN; TR; TT; TZ; UA; UG; US; UZ; VC; VN; YU; ZA; ZM; ZW
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  13/3, K/1 (Item 1 from file: 399) Links
CA SEARCH(R)
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143109795
                           CA: 143(7)109795k
                                                                     PATENT
Cathepsin Z inhibitors for treatment of rheumatoid arthritis and other autoimmune
Inventor (Author): Saltzman, Alan G.; Tang, Zhihua; Palejwala, Vaseem; Cavallo, Jean
Location: USA
Assignee: Aventis Pharmaceuticals Inc.
Patent: PCT International; WO 200565693 A2
                                                                              Date: 20050721
Application: WO 2004US41815 (20041214) *US 2003PV533330 (20031230)
Pages: 28 pp.
CODEN: PIXXD2
Language: English
Patent Classifications:
                 A61K-031/70A
Designated Countries: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY; BZ; CA;
CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; MA; MD; MG; MK; MN; MW; MX; NA; NI; NO; NZ; OM; PG; PH; PL; PT; RO; RU; SC; SD; SE; SG; SK; SL; SY; TJ; TM; TN; TR; TT; TZ; UA; UG; US; UZ; VC; VN; YU; ZA; ZM; ZW Designated Regional: BW; GH; GM; KE; LS; MW; MZ; NA; SD; SL; SZ; TZ; UG; ZM; ZW; AM; AZ; BY; KG; KZ; MD; RU; TJ; TM; AT; BE; BG; CH; CY; CZ; DE; DK; EE; ES; FI; FR; GB; GR; HU; IE; IS; IT; LT; LU; MC; NL; PL; PT; RO; SE; SI; SK; TR; BF; BJ; CF; CG; CI; CM: GA: GN: GO: GW: MI: MR: NE: SN: TD: TG
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CM; GA; GN; GQ; GW; ML; MR; NE; SN; TD; TG

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